

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
13 December 2001 (13.12.2001)

PCT

(10) International Publication Number
WO 01/94319 A1(51) International Patent Classification⁷: **C07D 237/14, 237/22, 237/32, 405/04, A61K 31/50, 31/501, 31/502, A61P 11/06**

3, NL-2253 VM Voorschoten (NL). CHRISTIAANS, Johannes; Zevenwouden 233, NL-3524 CR Utrecht (NL). BRUNDEL, Paulus; Matthias Withoosstraat 20, NL-3812 ST Amersfoort (NL).

(21) International Application Number: PCT/EP01/06230

(72) Inventor; and

(22) International Filing Date: 1 June 2001 (01.06.2001)

(75) Inventor/Applicant (for US only): STERK, Geert [NL/NL]; Stadhouderslaan 38, NL-3583 JJ Utrecht (NL).

(25) Filing Language: English

(74) Agent: RUPP, Herbert; c/o Byk Gulden Lomberg Chemische Fabrik GmbH, Byk-Gulden-Strasse 2, 78467 Konstanz (DE).

(30) Priority Data: 00111795.1 5 June 2000 (05.06.2000) EP

(81) Designated States (national): AE, AL, AU, BA, BG, BR, CA, CN, CO, CU, CZ, EC, EE, GE, HR, HU, ID, IL, IN, IS, JP, KR, LT, LV, MK, MX, NO, NZ, PL, RO, SG, SI, SK, UA, US, VN, YU, ZA, ZW.

(71) Applicants (for all designated States except US): BYK GULDEN LOMBERG CHEMISCHE FABRIK GMBH [DE/DE]; Patentabteilung, Byk-Gulden-Strasse 2, 78467 Konstanz (DE). BYK NEDERLAND BV [NL/NL]; Weerenweg 29, NL-1161 AG Zwanenburg (NL).

(84) Designated States (regional): Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR).

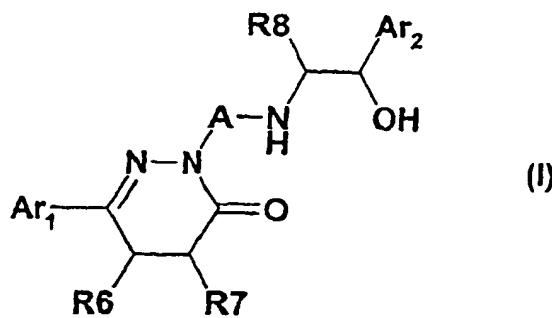
Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(72) Inventors (for all designated States except CA, US): HATZELMANN, Armin; Alter Wall 3, 78467 Konstanz (DE). BUNDSCHEUH, Daniela; Rheingutstrasse 17, 78462 Konstanz (DE). ELTZE, Manfrid; Schützenstrasse 20, 78462 Konstanz (DE). VAN DER LAAN, Yvonne; M. Schaftstraat 73, NL-2264 DP Leidschendam (NL). TIMMERMANN, Hendrik; De Savornin Lohmanplantsoen

(54) Title: COMPOUNDS EFFECTIVE AS BETA-2-ADRENORECEPTOR AGONISTS AS WELL AS PDE4-INHIBITORS

(57) Abstract: The compounds of formula I in which Ar₁, A, R₆, R₇, R₈ and Ar₂ have the meanings as given in the description are novel effective bronchial therapeutics.

Compounds effective as β 2-Adrenoreceptor Agonists as well as PDE4-Inhibitors

Field of application of the invention

The invention relates to novel compounds, which act simultaneously as β 2-adrenoreceptor agonist and as PDE4-inhibitor. They are used in the pharmaceutical industry for the production of medicaments.

Known technical background

International Patent Applications WO98/31674, WO99/31071 and WO99/31090 disclose phthalazinone derivatives having selective PDE4 inhibitory properties. In the International Patent Application WO94/12461 and in the European Patent Application EP 0 763 534 3-aryl-pyridazin-6-one respectively arylalkyl-diazinone derivatives are described as selective PDE4 inhibitors.

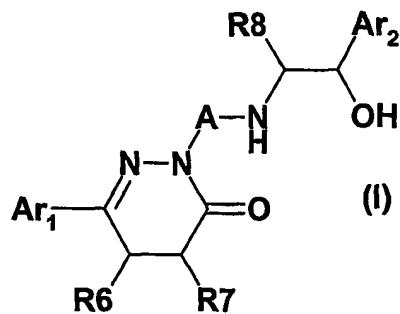
In the International Patent Application WO98/37894 the combined application of PDE-inhibitors with adenylat-cyclase-agonists is described. In this context β -sympathomimetics are mentioned as one possible example of an adenylat-cyclase-agonist.

In the International Patent Application WO00/12078 is disclosed the combined application of a PDE4 inhibitor with a beta adrenergic bronchodilator for the treatment of asthma and COPD.

Description of the invention

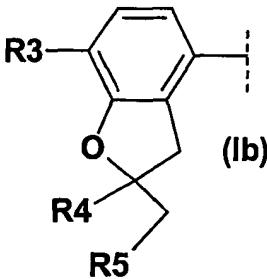
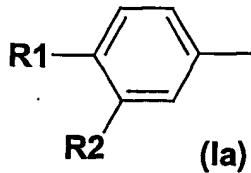
It has now been found that the pyridazinones, which are described in greater details below, have surprising and particularly advantageous properties.

The invention thus relates to compounds of the formula I



in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

R1 is hydroxyl, halogen, 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

R2 is hydroxyl, halogen, 1-8C-alkoxy, 3-7C-cycloalkoxy, 3-7C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

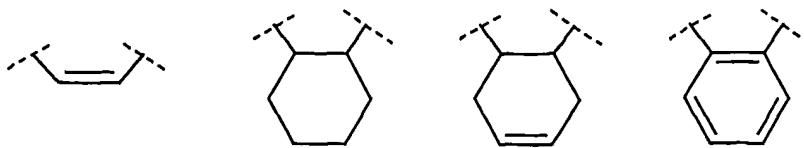
R4 is 1-4C-alkyl and

R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked 5-, 6- or 7-membered hydrocarbon ring, optionally interrupted by an oxygen or sulphur atom,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$ or $-Y-X-C_mH_{2m}-Z-C_nH_{2n}-$, wherein

X represents a bond, $-O-$ (oxygen), $-S-$ (sulfur), $-NH-$, $-C(O)-$, $-S(O)_2-$, $-C(O)-NH-$, $-NH-C(O)-$, $-C(S)-NH-$, $-NH-C(S)-$, $-NH-C(O)-NH-$ or $-NH-C(S)-NH-$,

Y represents a bond, phenylene, 4-8C-cycloalkylene or azacycloalkylene,

Z represents $-O-$, $-S-$, $-S(O)_2-$, $-NH-C(O)-$, $-C(O)-NH-$, $-NH-C(S)-$ or $-C(S)-NH-$,

m is an integer from 0 to 4,

n is an integer from 1 to 4,

R8 is hydrogen or 1-4C-alkyl,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11,

wherein

R9 is hydrogen, halogen, hydroxyl, amino, ureido $[-NH-C(O)-NH_2]$, formylamino $[-NH-C(O)H]$, 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy $[-O-C(O)-C_6H_4-CH_3]$, hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,

R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy $[-O-C(O)-C_6H_4-CH_3]$, hydroxymethyl or 1-4C-alkylcarbonyloxy,

R11 is hydrogen or halogen,

and the salts of these compounds.

1-4C-alkyl is a straight-chain or branched alkyl radical having 1 to 4 carbon atoms. Examples are the butyl, isobutyl, sec-butyl, tert-butyl, propyl, isopropyl, ethyl and methyl radicals.

Halogen within the meaning of the invention is chlorine, bromine or fluorine.

1-4C-alkoxy is a radical which, in addition to the oxygen atom, contains a straight-chain or branched alkyl radical having 1 to 4 carbon atoms. Alkoxy radicals having 1 to 4 carbon atoms which may be mentioned in this context are, for example, the butoxy, isobutoxy, sec-butoxy, tert-butoxy, propoxy, iso-propoxy, ethoxy and methoxy radicals.

3-7C-cycloalkoxy stands for cyclopropyloxy, cyclobutyloxy, cyclopentyloxy, cyclohexyloxy and cycloheptyloxy.

3-7C-cycloalkylmethoxy stands for cyclopropylmethoxy, cyclobutylmethoxy, cyclopentylmethoxy, cyclohexylmethoxy and cycloheptylmethoxy.

1-4C-alkoxy which is completely or predominantly substituted by fluorine is, for example, the 2,2,3,3,3-pentafluoropropoxy, the perfluoroethoxy, the 1,2,2-trifluoroethoxy and in particular the 1,1,2,2-tetrafluoroethoxy, the 2,2,2-trifluoroethoxy, the trifluoromethoxy and the difluoromethoxy radical, of which the difluoromethoxy radical is preferred. "Predominantly" in this connection means that more than half of the hydrogen atoms of the 1-4C-alkoxy group are replaced by fluorine atoms.

As spiro-linked 5-, 6- or 7-membered hydrocarbon rings, optionally interrupted by an oxygen or sulphur atom, may be mentioned the cyclopentane, cyclohexane, cycloheptane, tetrahydrofuran, tetrahydropyran and the tetrahydrothiophen ring.

Possible radicals $-(CH_2)_m-$, in which m can have the meaning 0, 1, 2, 3 or 4 are straight chain or branched divalent alkylene radicals. Examples which may be mentioned are the butylene, isobutylene, sec-butylene, tert-butylene, propylene, isopropylene, ethylene and the methylene radical. The radical $-(CH_2)_m-$ represents a bond, if m is zero.

Possible radicals $-C_mH_{2m}-$, in which m can have the meaning 0, 1, 2, 3 or 4 are straight chain or branched divalent alkylene radicals. Examples which may be mentioned are the butylene, isobutylene, sec-butylene, tert-butylene, propylene, isopropylene, ethylene and the methylene radical. The radical $-C_mH_{2m}-$ represents a bond, if m is zero.

Possible radicals $-(CH_2)_n-$, in which n can have the meaning 1, 2, 3 or 4 are straight chain or branched divalent alkylene radicals. Examples which may be mentioned are the butylene, isobutylene, sec-butylene, tert-butylene, propylene, isopropylene, ethylene and the methylene radical.

Possible radicals $-C_nH_{2n}-$, in which n can have the meaning 1, 2, 3 or 4 are straight chain or branched divalent alkylene radicals. Examples which may be mentioned are the butylene, isobutylene, sec-butylene, tert-butylene, propylene, isopropylene, ethylene and the methylene radical.

An 1-4C-alkylcarbonylamino radical is, for example, the propionylamino $[C_3H_7C(O)NH-]$ and the acetyl-amino radical $[CH_3C(O)NH-]$.

4-8C-cycloalkylene stands for cyclobutylene, cyclopentylene, cyclohexylene, cycloheptylene or cyclooctylene. Preferred examples are 1,4-cyclohexylene and 1,3-cyclohexylene.

Azacycloalkylene stands for a divalent 5-7C-cycloalkylene radical, in which one or two of the ring carbon atoms are substituted by a nitrogen atom. Examples which may be mentioned are the 4,1-piperidinylene, the 1,4-piperidinylene, the 1,4-homopiperazinylene, the 1,4-piperazinylene and the 3,1-pyrrolidinylene radical.

Mono- or Di-1-4C-alkylamino radicals contain in addition to the nitrogen atom, one or two of the above-mentioned 1-4C-alkyl radicals. Examples which may be mentioned are the di-1-4C-alkylamino radicals, especially the dimethylamino, the diethylamino and the diisopropylamino radical.

Mono- or Di-1-4C-alkylaminocarbonyloxy radicals contain in addition to the carbonyl group one of the above-mentioned mono- or di-1-4C-alkylamino radicals. Examples which may be mentioned are the N-methyl-, the N,N-dimethyl-, the N-ethyl-, the N-propyl-, the N,N-diethyl- and the N-isopropylamino-carbonyloxy radical.

1-4C-Alkylcarbonyloxy stands for a carbonyloxy group to which one of the above-mentioned 1-4C-alkyl radicals is bonded. An example is the acetoxy radical $[\text{CH}_3\text{C}(\text{O})\text{O}]$.

1-4C-Alkylsulfonylamino is a sulfonylamino ($-\text{SO}_2\text{NH}-$) group to which one of the above-mentioned 1-4C-alkyl radicals is bonded. An example is the methanesulfonylamino radical $(\text{CH}_3\text{SO}_2\text{NH})$.

1-4C-Alkylsulfonylmethyl is a sulfonylmethyl ($-\text{SO}_2\text{CH}_2-$) group to which one of the above-mentioned 1-4C-alkyl radicals is bonded. An example is the methanesulfonylmethyl radical $(\text{CH}_3\text{SO}_2\text{CH}_2)$.

1-4C-Alkoxy-1-4C-alkyl stands for one of the above-mentioned 1-4C-alkyl radicals which is substituted by one of the above-mentioned 1-4C-alkoxy radicals. Examples which may be mentioned are the methoxymethyl, methoxyethyl and the butoxyethyl radical.

Ar_2 represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11. Examples which may be mentioned for the phenyl radical substituted by R9, R10 and R11 are phenyl, 4-amino-3-chloro-5-cyanophenyl, 4-hydroxy-3-hydroxymethylphenyl, 4-amino-3-cyanophenyl, 4-amino-3,5-dichlorophenyl, 4-amino-3-chloro-5-trifluoromethylphenyl, 4-aminophenyl, 3,4-dihydroxyphenyl, 3,5-dihydroxyphenyl, 3,5-bis-tolylcarbonyloxyphenyl, 4-hydroxy-3-ureido-phenyl, 3-formylamino-4-hydroxyphenyl, 4-hydroxyphenyl, 3-amino-5-hydroxyphenyl, 2-fluorophenyl, 3-amino-5-hydroxymethylphenyl, 3,5-di-tert-butylcarbonyloxyphenyl, 3,5-di-isopropylcarbonyloxyphenyl,

2-chloro-4-hydroxyphenyl, 2-fluoro-4-hydroxyphenyl, 3-fluoro-4-hydroxyphenyl, 4-hydroxy-3-methoxyphenyl, 4-hydroxy-3-methylsulfonamidophenyl and 4-hydroxy-3-methylsulfonylmethylphenyl.

Preferred examples of Ar_2 are 4-amino-3-chloro-5-cyanophenyl, 4-hydroxy-3-hydroxymethylphenyl, 4-amino-3-cyanophenyl, 4-amino-3-chloro-5-trifluoromethylphenyl and 4-amino-3,5-dichlorophenyl.

Suitable salts for compounds of the formula I - depending on substitution - are all acid addition salts or all salts with bases. Particular mention may be made of the pharmacologically tolerable inorganic and organic acids and bases customarily used in pharmacy. Those suitable are, on the one hand, water-soluble and water-insoluble acid addition salts with acids such as, for example, hydrochloric acid, hydrobromic acid, phosphoric acid, nitric acid, sulphuric acid, acetic acid, citric acid, D-gluconic acid, benzoic acid, 2-(4-hydroxybenzoyl)benzoic acid, butyric acid, sulphosalicylic acid, maleic acid, lauric acid, malic acid, fumaric acid, succinic acid, oxalic acid, tartaric acid, embonic acid, stearic acid, toluenesulphonic acid, methanesulphonic acid or 3-hydroxy-2-naphthoic acid, the acids being employed in salt preparation - depending on whether a mono- or polybasic acid is concerned and depending on which salt is desired - in an equimolar quantitative ratio or one differing therefrom.

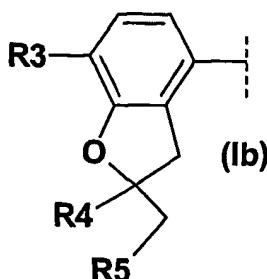
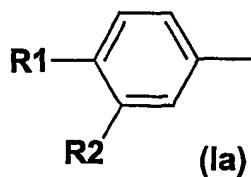
On the other hand, salts with bases are - depending on substitution - also suitable. As examples of salts with bases are mentioned the lithium, sodium, potassium, calcium, aluminium, magnesium, titanium, ammonium, meglumine or guanidinium salts, here, too, the bases being employed in salt preparation in an equimolar quantitative ratio or one differing therefrom.

Pharmacologically intolerable salts, which can be obtained, for example, as process products during the preparation of the compounds according to the invention on an industrial scale, are converted into pharmacologically tolerable salts by processes known to the person skilled in the art.

According to expert's knowledge the compounds of the invention as well as their salts may contain, e.g. when isolated in crystalline form, varying amounts of solvents. Included within the scope of the invention are therefore all solvates and in particular all hydrates of the compounds of formula I as well as all solvates and in particular all hydrates of the salts of the compounds of formula I.

Compounds of the formula I to be emphasized are those in which

Ar_1 represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



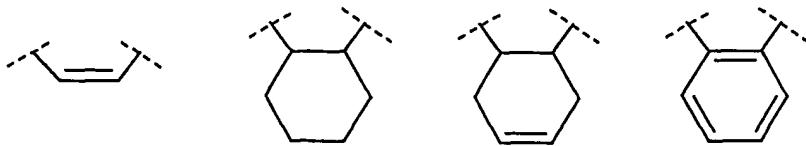
wherein

- R1 is 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is 1-4C-alkyl and
- R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane, cyclohexane, tetrahydrofuran or tetrahydropyran ring,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



and A either represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein either

- X represents a bond,
- Y represents a bond,
- m is an integer from 1 to 4, and
- n is an integer from 1 to 4,
- or
- X represents a bond, -O-, -S-, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,
- Y represents 1,4-phenylene, 1,3-phenylene, 1,4-cyclohexylene, 1,3-cyclohexylene or 1,3-cyclopentylene,
- m is an integer from 0 to 4, and

n is an integer from 1 to 4,
 or
 X represents a bond, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,
 Y represents 4,1-piperidinylene,
 m is 0, and
 n is an integer from 1 to 4,
 or
 X represents a bond, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,
 Y represents 1,4-piperazinylene,
 m is an integer from 1 to 4, and
 n is an integer from 1 to 4,

or

A represents -Y-X-C_mH_{2m}-Z-C_nH_{2n}-, wherein

X represents a bond, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,
 Y represents 4,1-piperidinylene,
 Z represents a bond, -O-, -S-, -S(O)₂- or -C(O)-NH-,
 m is an integer from 1 to 4, and
 n is an integer from 1 to 4,
 or
 X represents a bond, -O-, -S-, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,
 Y represents 1,4-phenylene, 1,3-phenylene, 1,4-cyclohexylene, 1,3-cyclohexylene or
 1,3-cyclopentylene,
 Z represents a bond, -O-, -S-, -S(O)₂- or -C(O)-NH-,
 m is an integer from 1 to 4, and
 n is an integer from 1 to 4,

R8 is hydrogen, methyl or ethyl,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein

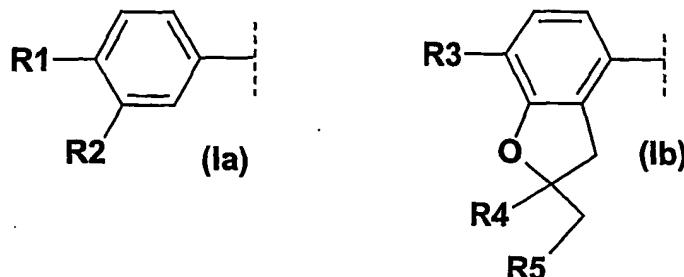
R9 is hydrogen, halogen, hydroxyl, amino, ureido [-NH-C(O)-NH₂], formylamino [-NH-C(O)H], 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,

R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl or 1-4C-alkylcarbonyloxy,

R11 is hydrogen or halogen,

and the salts of these compounds.

Compounds of the formula I which are particularly to be emphasized are those in which Ar₁ represents a phenyl radical of formula (Ia) substituted by R₁ and R₂ or a dihydrobenzofuranyl radical of formula (Ib) substituted by R₃, R₄ and R₅



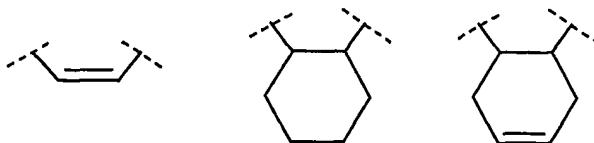
wherein

- R1 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is 1-2C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is methyl and
- R5 is hydrogen,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane or cyclohexane ring.

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



and A either represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein

either

X represents a bond,

Y represents a bond,

m is an integer from 1 to 4, and

n is an integer from 1 to 4.

or

X represents a bond, -O- or -C(O)-NH-

Y represents 1,4-phenylene or 1,4-cyclohexylene

m is an integer from 0 to 4, and
n is an integer from 1 to 4,
or
X represents a bond, -C(O)-, -S(O)₂- or -C(S)-NH-,
Y represents 4,1-piperidinylene,
m is 0, and
n is an integer from 1 to 4,

or

A represents -Y-X-C_mH_{2m}-Z-C_nH_{2n}-, wherein

either
X represents a bond or -C(O)-,
Y represents 4,1-piperidinylene,
Z represents a bond, -S- or -S(O)₂-,
m is an integer from 1 to 4, and
n is an integer from 1 to 4,
or
X represents a bond, -O- or -C(O)-NH-,
Y represents 1,4-phenylene or 1,4-cyclohexylene,
Z represents a bond,
m is an integer from 1 to 4, and
n is an integer from 1 to 4,

R8 is hydrogen,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein

R9 is hydrogen, hydroxyl or amino,

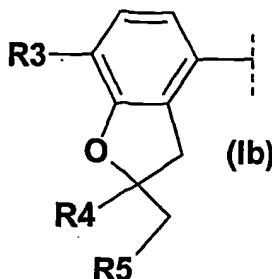
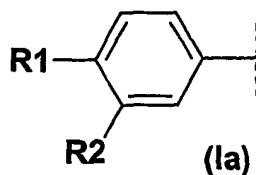
R10 is hydrogen, halogen, cyano, trifluoromethyl or hydroxymethyl,

R11 is hydrogen or halogen,

and the salts of these compounds.

Preferred compounds of formula I are those in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

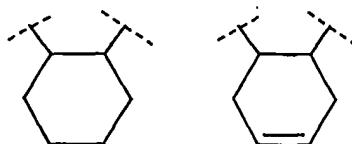
R1 is methoxy or ethoxy,

R2 is methoxy or ethoxy,

R3 is methoxy,

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane ring,

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



and A either represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein

either

X represents a bond,

Y represents a bond,

m is an integer from 1 to 4, and

n is an integer from 1 to 4,

or

X represents a bond, -O- or -C(O)-NH-,

Y represents 1,4-phenylene,

m is an integer from 0 to 1, and

n is an integer from 1 to 4,

or

X represents a bond, -C(O)-, -S(O)₂- or -C(S)-NH-,

Y represents 4,1-piperidinylene,

m is 0, and

n is an integer from 1 to 4,

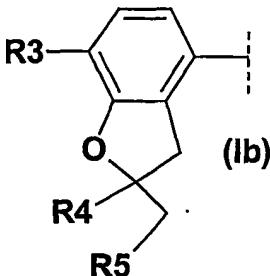
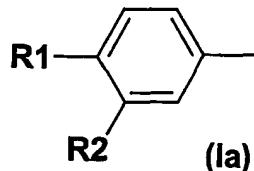
or

A represents $-Y-X-C_mH_{2m}-Z-C_nH_{2n}-$, wherein

- X represents a bond or $-C(O)-$,
- Y represents 4,1-piperidinylene,
- Z represents a bond, $-S-$ or $-S(O)_2-$,
- m is 2 or 3, and
- n is 2 or 3,
- R8 is hydrogen,
- Ar₂ is phenyl, 4-amino-3-chloro-5-cyanophenyl, 4-amino-3-chloro-5-trifluoromethylphenyl, 4-hydroxy-3-hydroxymethylphenyl, 4-amino-3-cyanophenyl or 4-amino-3,5-dichlorophenyl, and the salts of these compounds.

One embodiment (embodiment A) of the compounds of formula I are those, in which

- Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



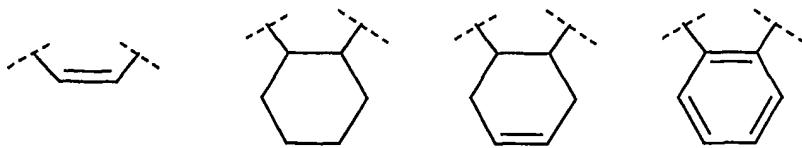
wherein

- R1 is hydroxyl, halogen, 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is hydroxyl, halogen, 1-8C-alkoxy, 3-7C-cycloalkoxy, 3-7C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is 1-4C-alkyl and
- R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked 5-, 6- or 7-membered hydrocarbon ring, optionally interrupted by an oxygen or sulphur atom,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents $-(CH_2)_m-Y-X-(CH_2)_n-$, wherein

- X represents a bond, -O- (oxygen), -S- (sulfur), -NH-, -C(O)-, -C(O)-NH- or -NH-C(O)-NH-,
- Y represents a bond, phenylene, 4-8C-cycloalkylene or azacycloalkylene,
- m is an integer from 0 to 4,
- n is an integer from 1 to 4,

R8 is hydrogen or 1-4C-alkyl,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11,

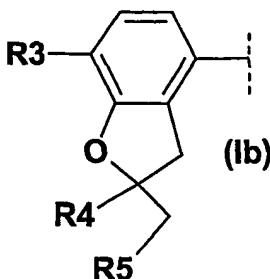
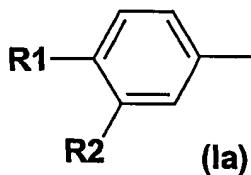
wherein

- R9 is hydrogen, halogen, hydroxyl, amino, ureido [-NH-C(O)-NH₂], formylamino [-NH-C(O)H], 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,
- R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl or 1-4C-alkylcarbonyloxy,
- R11 is hydrogen or halogen,

and the salts of these compounds.

Compounds of the formula I of embodiment A to be emphasized are those in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



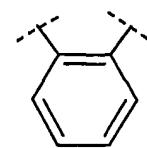
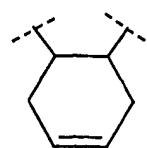
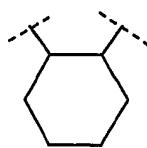
wherein

- R1 is 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is 1-4C-alkyl and
- R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane, cyclohexane, tetrahydrofuran or tetrahydropyran ring,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



- A represents $-(CH_2)_m-Y-X-(CH_2)_n-$, wherein
 - X represents a bond, -O- (oxygen), -NH-, -C(O)- or -C(O)-NH-,
 - Y represents a bond, phenylene, cyclopentylene, cyclohexylene, piperidinylene or piperazinylene,
 - m is an integer from 0 to 4,
 - n is an integer from 1 to 4,
- R8 is hydrogen, methyl or ethyl,
- Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein

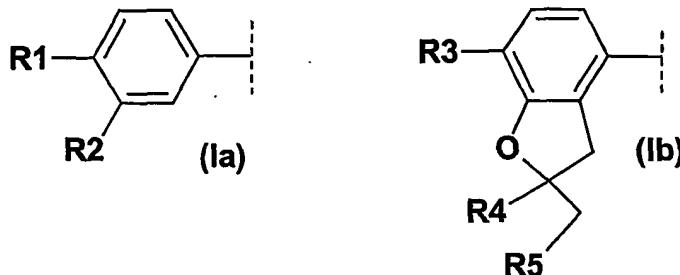
R9 is hydrogen, halogen, hydroxyl, amino, ureido [-NH-C(O)-NH₂], formylamino [-NH-C(O)H], 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,

R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl or 1-4C-alkylcarbonyloxy,

R11 is hydrogen or halogen,
and the salts of these compounds.

Compounds of the formula I of embodiment A which are particularly to be emphasized are those in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



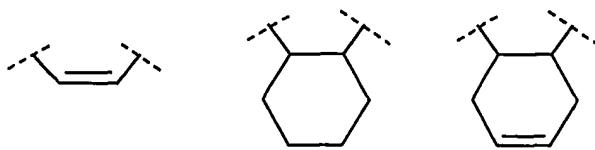
wherein

R1 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
 R2 is 1-2C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
 R3 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
 R4 is methyl and
 R5 is hydrogen,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane or cyclohexane ring,

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents $-(CH_2)_m-Y-X-(CH_2)_n-$, wherein

X represents a bond or $-C(O)-$,

Y represents a bond, 1,4-phenylene, 1,4-cyclohexylene or 4,1-piperidinylene,

m is an integer from 0 to 4,

n is an integer from 1 to 4,

R8 is hydrogen,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein

R9 is hydrogen, hydroxyl or amino,

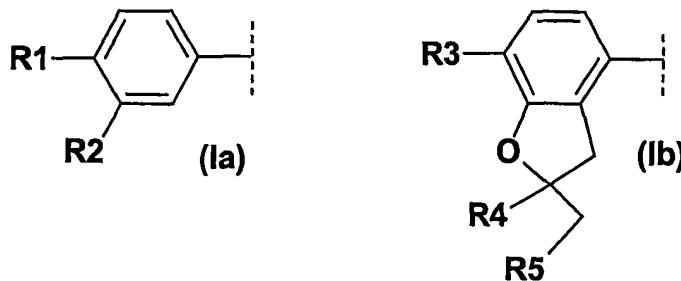
R10 is hydrogen, halogen, cyano, trifluoromethyl or hydroxymethyl,

R11 is hydrogen or halogen,

and the salts of these compounds.

Preferred compounds of formula I of embodiment A are those in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

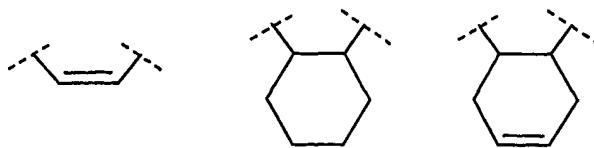
R1 is methoxy, ethoxy or difluoromethoxy,

R2 is methoxy or ethoxy,

R3 is methoxy,

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane ring,

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents $-(CH_2)_m-Y-X-(CH_2)_n-$, wherein

X represents a bond,

Y represents a bond or 1,4-phenylene,

m is an integer from 0 to 4,

n is an integer from 1 to 4,

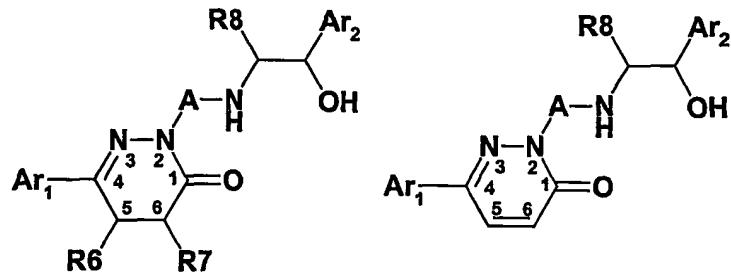
R8 is hydrogen,

Ar₂ is phenyl, 4-amino-3-chloro-5-cyanophenyl, 4-amino-3-chloro-5-trifluoromethylphenyl, 4-hydroxy-3-hydroxymethylphenyl, 4-amino-3-cyanophenyl or 4-amino-3,5-dichlorophenyl,

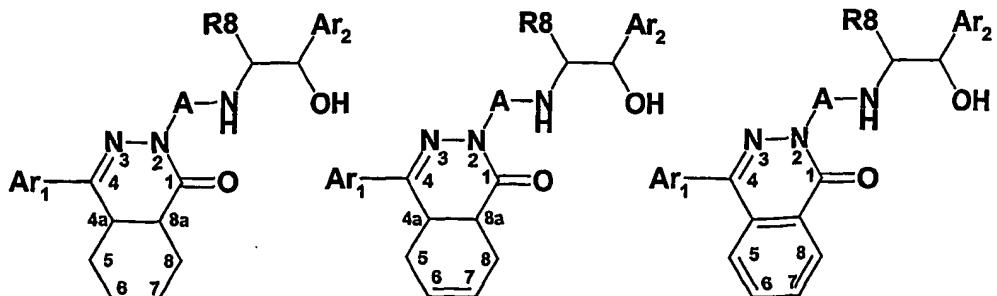
and the salts of these compounds.

For the purpose of this invention the following numbering is used in the phthalazinone respectively pyridazinone part of the compounds of formula I:

Numbering pyridazinone compounds

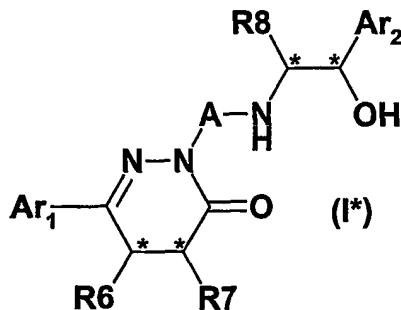


Numbering phthalazinone compounds



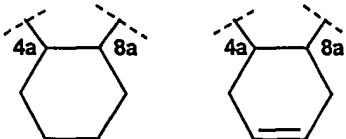
The compounds of formula I are chiral compounds with - depending of the meaning of Ar₁ - a chiral center in the dihydrobenzofuranyl radical, if the substituents -R4 and -CH₂R5 are not identical. However, preferred are those compounds, in which the substituents -R4 and -CH₂R5 are identical or together and with inclusion of the carbon atoms to which they are bonded form a spiro-connected 5-, 6- or 7-membered hydrocarbon ring.

Further possible chiral centers in the compounds of formula I are marked in the following formula I* with an asterix (*):



The invention includes all conceivable pure diastereomers and pure enantiomers, as well as all mixtures thereof independent from the ratio, including the racemates.

In those cases, wherein R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



those compounds are preferred, in which the hydrogen atoms in the positions 4a and 8a are cis-configured. Especially preferred in this connection are those compounds, in which the absolute configuration (according to the rules of Cahn, Ingold and Prelog) is S in the position 4a and R in the position 8a.

(4a,8a)-cis-Racemates can be split up into the corresponding enantiomers by methods known by a person skilled in the art. Preferably the racemic mixtures are separated into two diastereomers during the preparation with the help of an optical active separation agent on the stage of the cyclohexane-carboxylic acids or the 1,2,3,6-tetrahydrobenzoic acids. As separation agents may be mentioned, for example, optical active amines such as the (+)- and (-)-forms of 1-phenylethylamine [(R)-(+)-1-phen-

ylethylamine = D- α -methylbenzylamine or (S)-(-)-1-phenylethylamine = L- α -methylbenzylamine) and ephedrine, the optical active alkaloids quinine, cinchonine, cinchonidine and brucine.

The preparation of (4aS, 8aR) configurated 4-(3,4-dialkoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-ones or 4-(3,4-dialkoxyphenyl)-4a,5,6,7,8,8a-hexahydro-2H-phthalazin-1-ones is described, for example, in the international application WO98/31674.

The preparation of (4aS, 8aR) configurated 4-(2,3-dihydro-7-alkoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-4a,5,6,7,8,8a-hexahydro-2H-phthalazin-1-ones is described, for example, in the international application WO99/31090.

One group of preferred compounds of formula I are those in which the divalent radical A has a longitudinal extension of about 4 to about 6 carbon carbon bonds. Examples which may be mentioned in this connection are tetramethylene, pentamethylene, hexamethylene or phenylethylene.

Another group of preferred compounds of formula I are those in which Y represents a 1,4-phenylene or 4,1-piperidinylene radical.

The invention further relates to processes for the preparation of compounds of formula I and their salts.

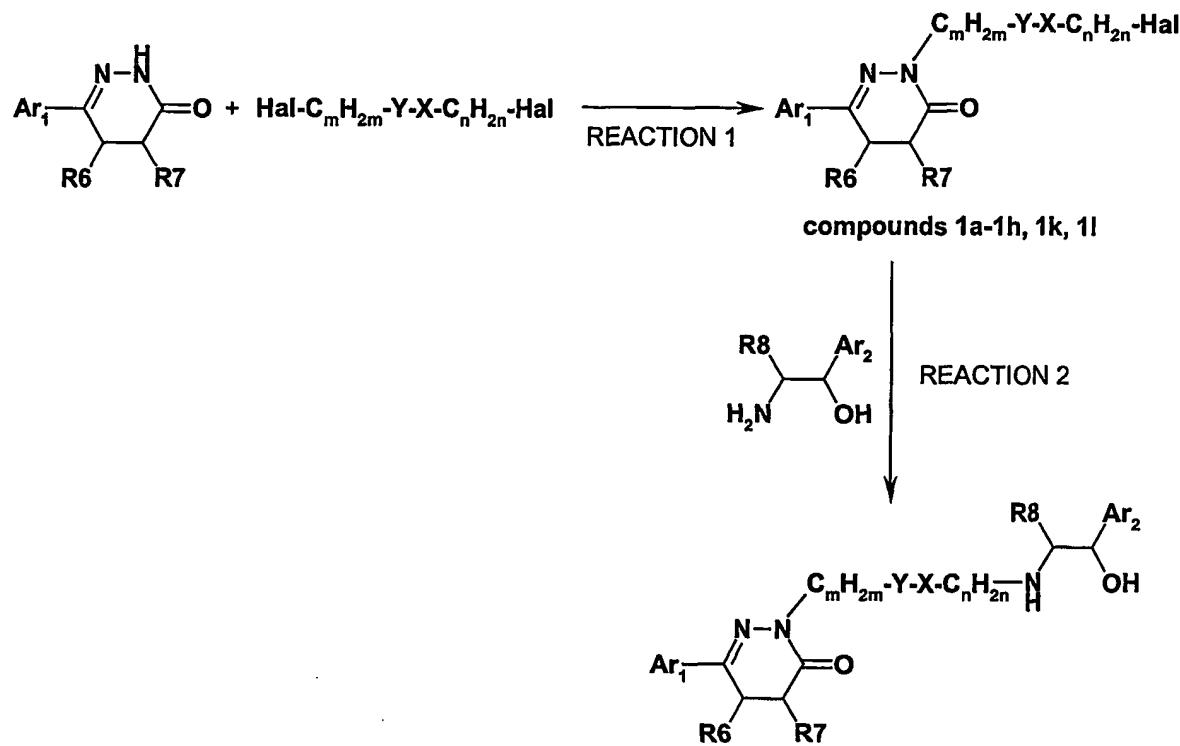
Several alternative synthetic routes to compounds of formula I are described in the Reaction Schemes I to IX.

Reaction Scheme I describes the preparation of compounds of formula I starting from 2H-phthalazin-1-one or 2H-pyridazin-1-one compounds. These are reacted in a first reaction with appropriate α,ω -dihalogenalkanes (if A represents an alkylene chain) to give the corresponding 2-(ω -halogenalkyl)phthalazin-1-ones or 2-(ω -halogenalkyl)pyridazinones. In a second reaction these compounds are then converted to compounds of formula I through reaction with an 2-amino-1-(phenyl)ethanol derivative. Compounds of formula I in which A contains additional to the alkylene chain further groups, for example a 1,4-phenylene radical, can be prepared in an analogous manner.

Reaction Scheme I:

[particularly suitable for compounds of formula I, in which

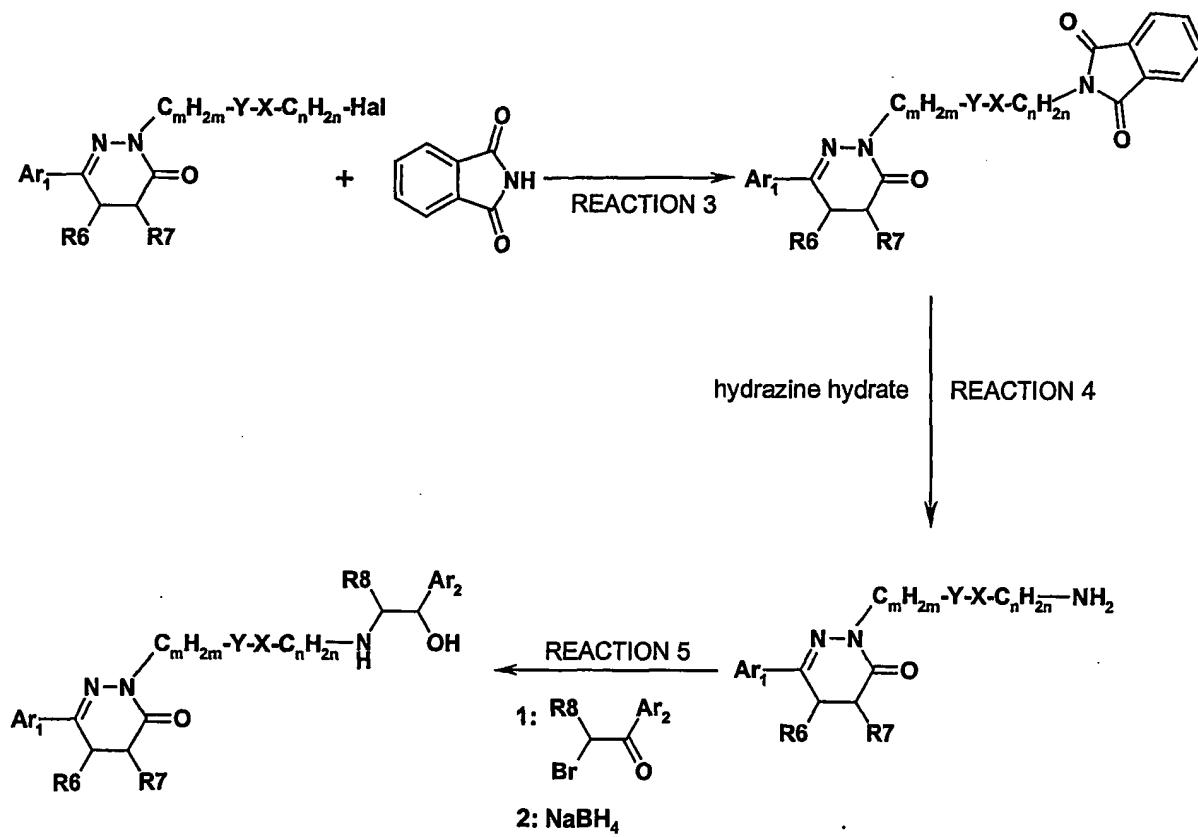
A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$ wherein
 X represents a bond,
 Y represents a bond, phenylene or 4-8C-cycloalkylene,
 m is an integer from 0 to 4,
 n is an integer from 1 to 4]



Reaction Scheme II:

[particularly suitable for compounds of formula I, in which

- A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$ wherein
- X represents a bond,
- Y represents a bond, phenylene or 4-8C-cycloalkylene,
- m is an integer from 0 to 4,
- n is an integer from 1 to 4]



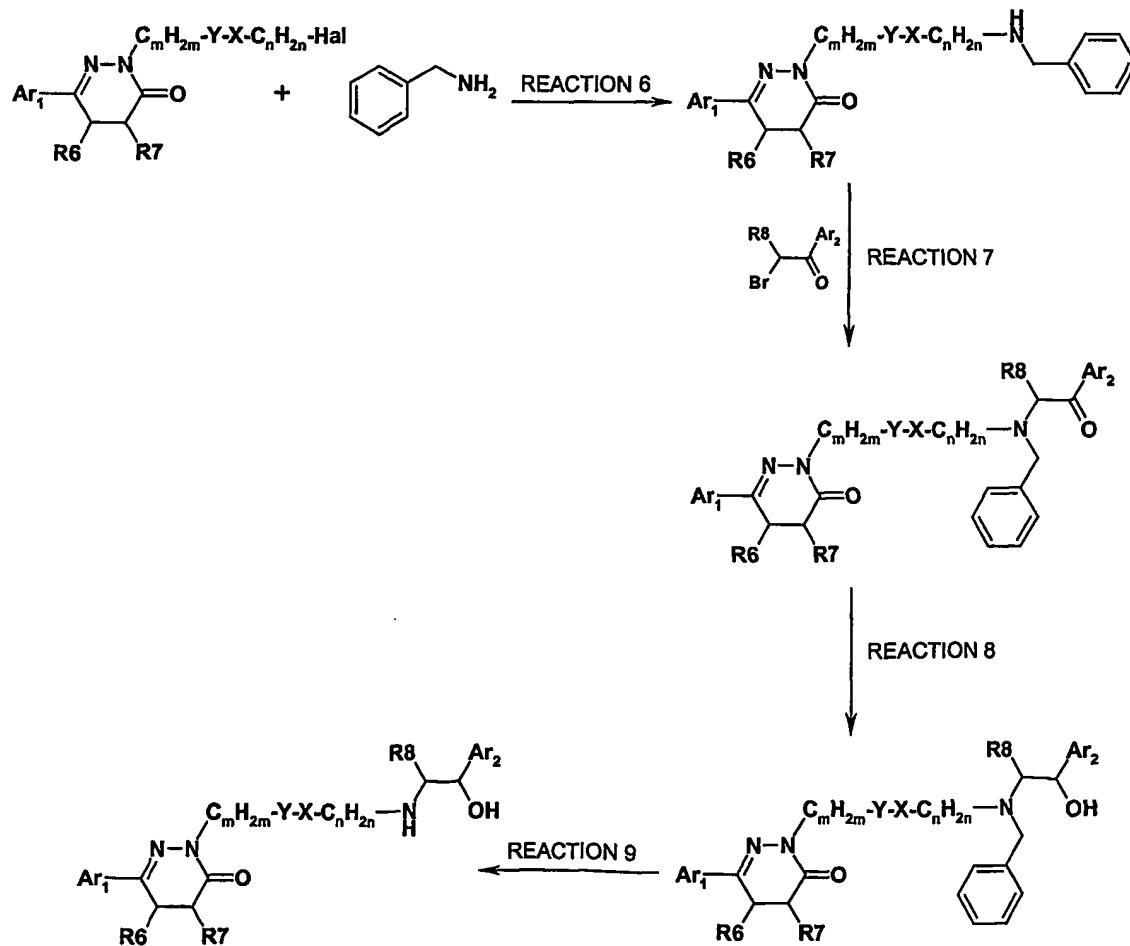
Reaction Scheme II starts with the 2-(ω -halogenalkyl)phthalazin-1-ones or 2-(ω -halogenalkyl)pyridazinones already described in Reaction Scheme I. In a first reaction the halogen atom is replaced by an amino group via a Gabriel synthesis (for example as described in Angew. Chem. 80, 986-996). The 1-(phenyl)ethanol part of the compounds of formula I is introduced with the help of a 2-bromoacetophenone derivative followed by a reduction step.

Reaction Scheme III also starts with the 2-(ω -halogenalkyl)phthalazin-1-ones or 2-(ω -halogenalkyl)pyridazinones already described in Reaction Scheme I. In this alternative the amino function is introduced through reaction with benzylamine. The 1-(phenyl)ethanol part of the compounds of formula I is again introduced with the help of a 2-bromoacetophenone derivative followed by a reduction step. Finally the benzyl group is removed, for example, by catalytic hydrogenation.

Reaction Scheme III:

[particularly suitable for compounds of formula I, in which

- A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$ wherein
- X represents a bond,
- Y represents a bond, phenylene or 4-8C-cycloalkylene,
- m is an integer from 0 to 4,
- n is an integer from 1 to 4]

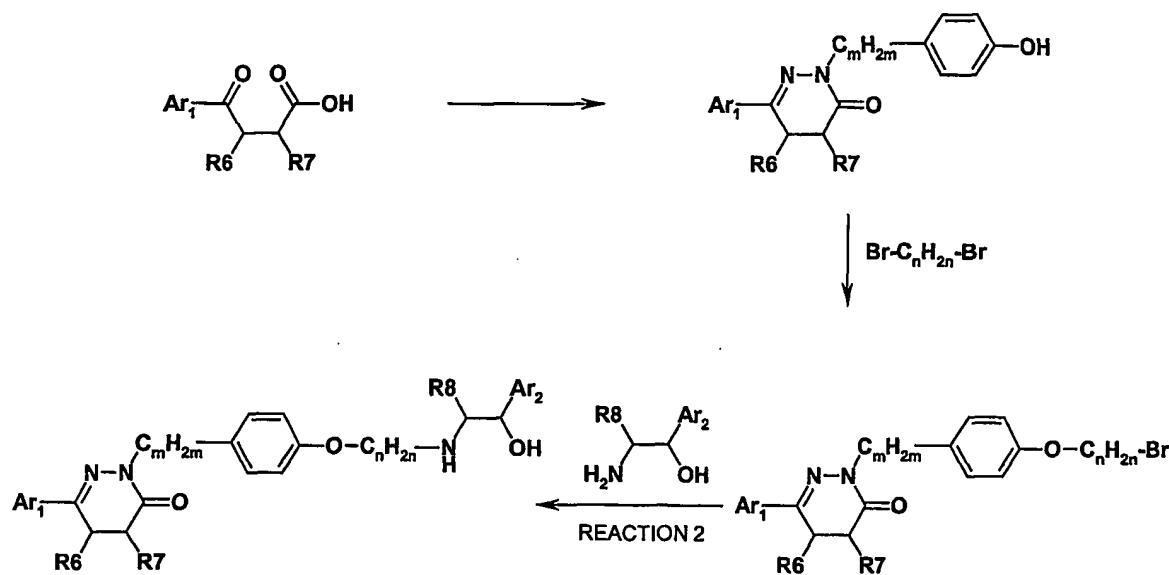


Reaction scheme IV:

[particularly suitable for compounds of formula I, in which

A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein
 X represents -O- (oxygen) or -S- (sulfur),
 Y represents phenylene or 4-8C-cycloalkylene,
 m is an integer from 0 to 4,
 n is an integer from 1 to 4];

Exemplarily shown is X = -O- and Y = phenylene:



Reaction scheme V:

[particularly suitable for compounds of formula I, in which

A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein

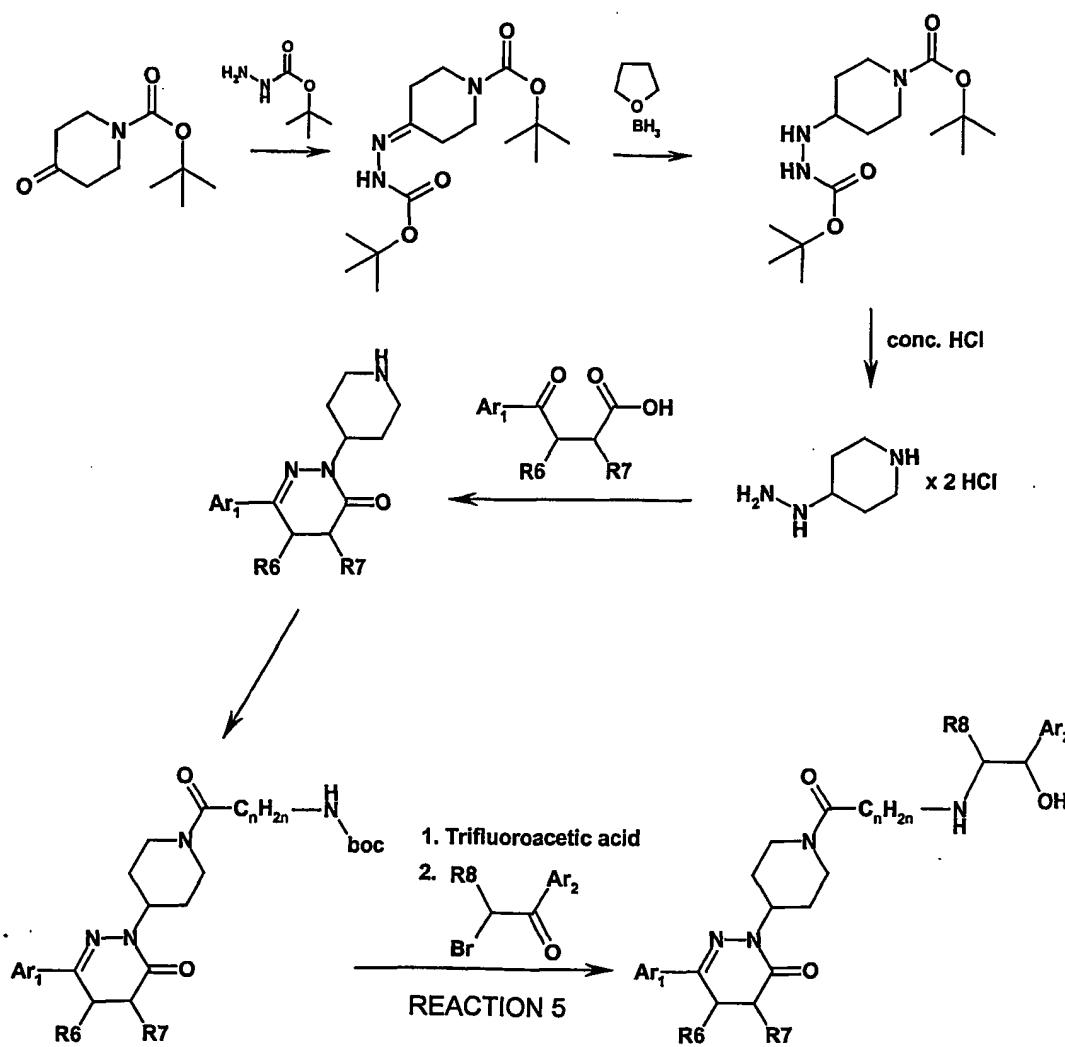
X represents $-C(O)-$ or $-S(O)_2-$,

Y represents azacycloalkylene,

m is 0,

n is an integer from 1 to 4];

Exemplarily shown is X = $-C(O)-$ and Y = 4,1-piperidylene:



Reaction scheme VI:

[particularly suitable for compounds of formula I, in which

A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein

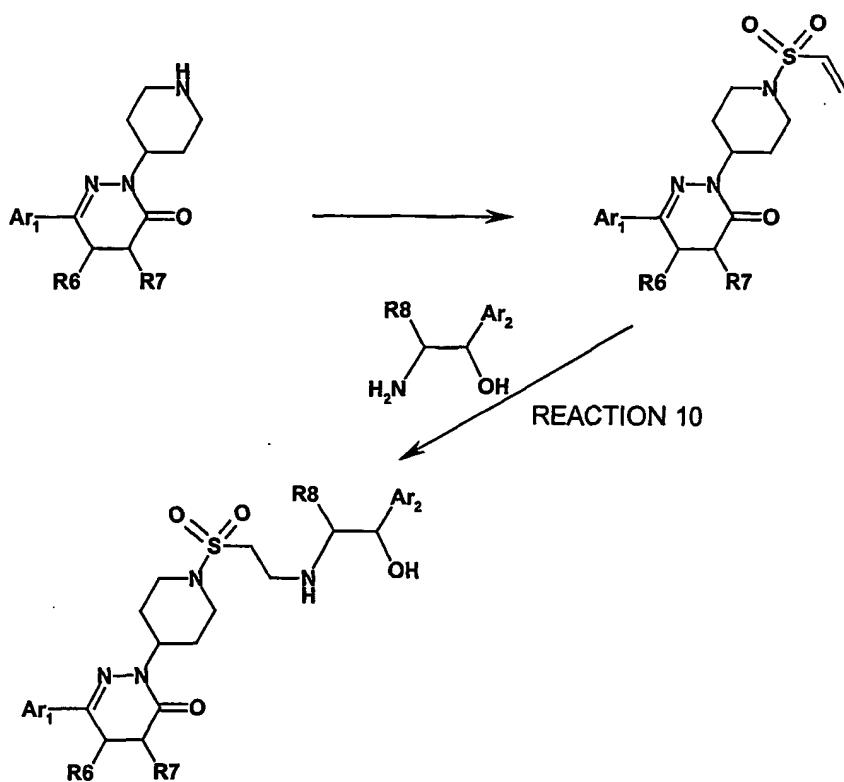
X represents $-S(O)_2-$,

Y represents azacycloalkylene,

m is 0,

n is 2]

Exemplarily shown is Y = 4,1-piperidylene:



Reaction scheme VII:

[particularly suitable for compounds of formula I, in which

A represents $-Y-X-C_mH_{2m}-Z-C_nH_{2n}-$, wherein

X represents a bond,

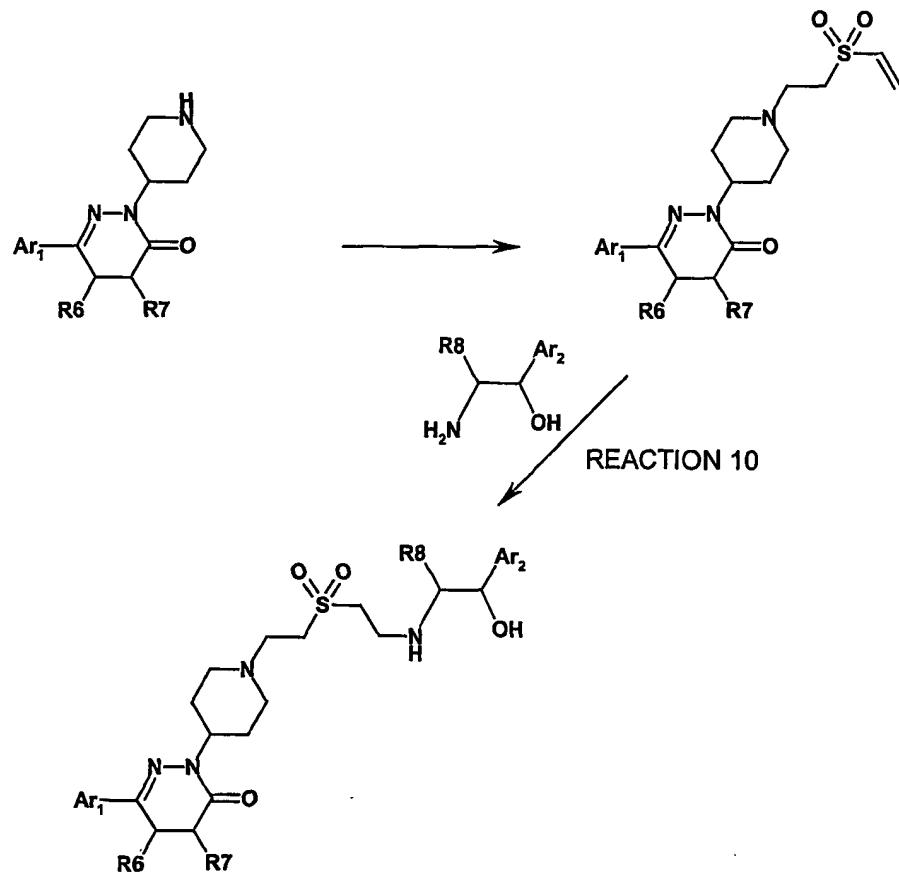
Y represents azacycloalkylene,

Z represents $-S(O)_2-$,

m is an integer from 1 to 4,

n is 2];

Exemplarily shown is Y = 4,1-piperidylene:



Reaction scheme VIII:

[particularly suitable for compounds of formula I, in which

A represents $-Y-X-C_mH_{2m}-Z-C_nH_{2n}-$, wherein

X represents $-C(O)-$,

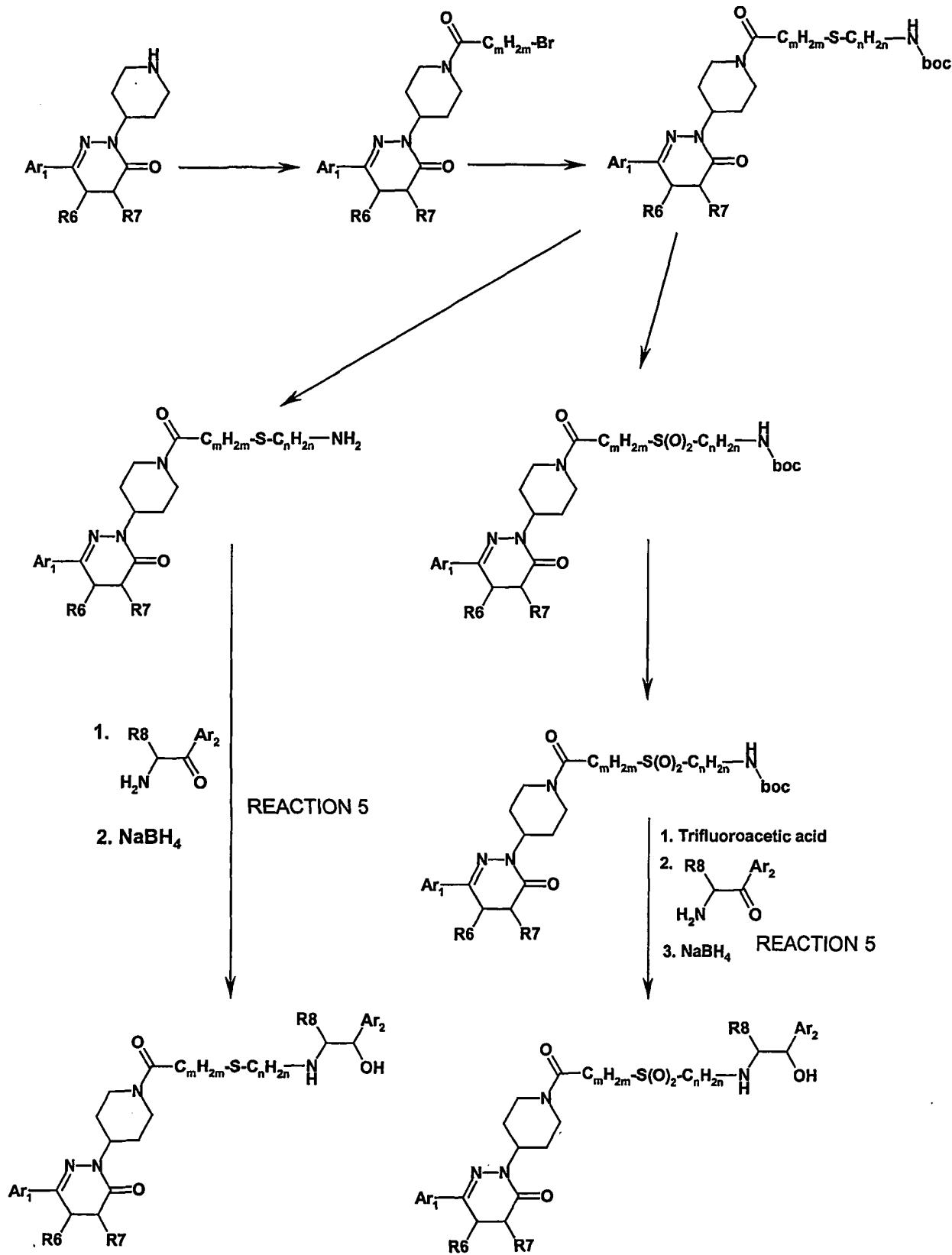
Y represents azacycloalkylene,

Z represents $-O-$, $-S-$ or $-S(O)_{2-}$,

m is an integer from 1 to 4,

n is an integer from 1 to 4];

Exemplarily shown is Y = 4,1-piperidylene and Z = -S- or -S(O)₂-:

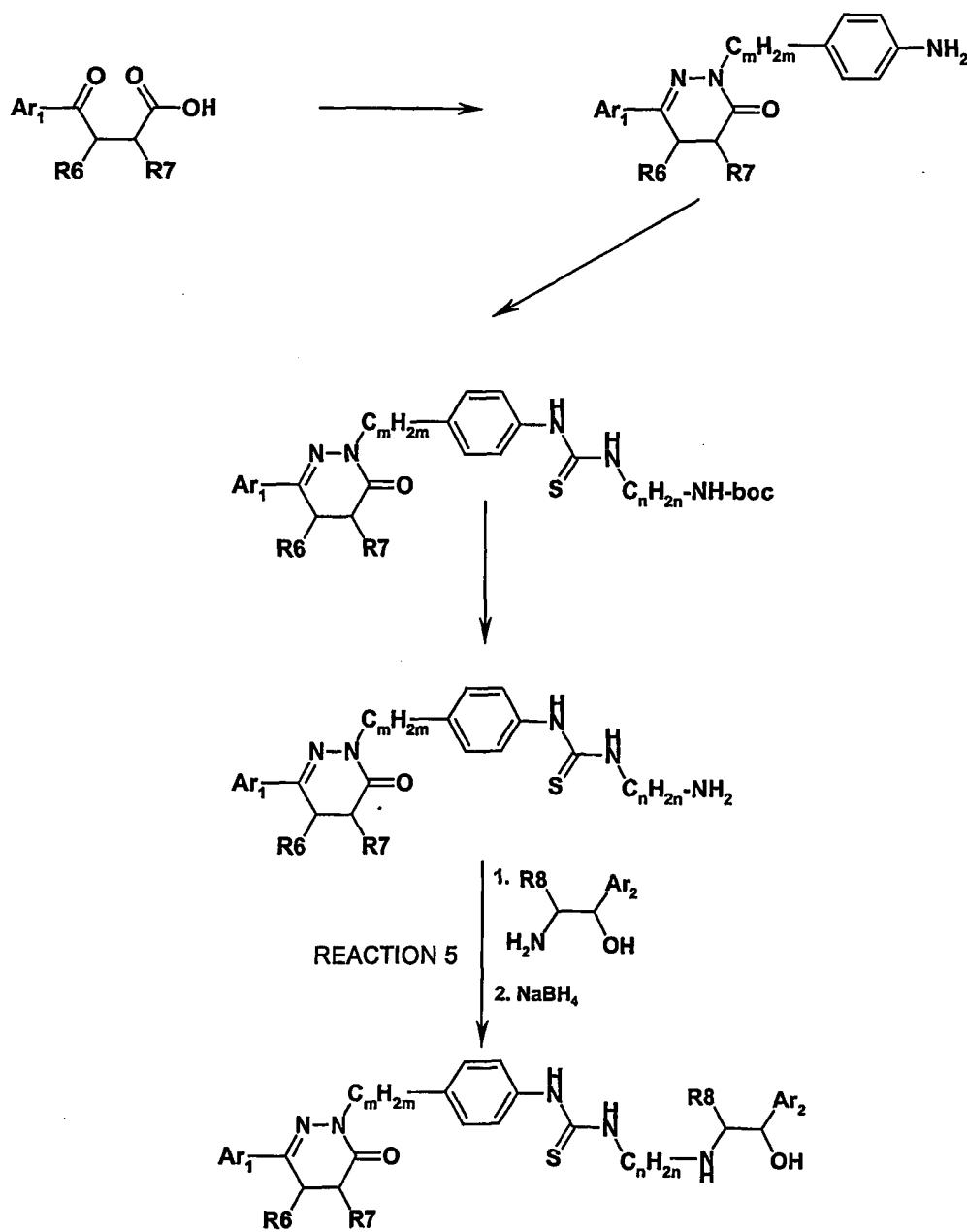


Reaction scheme IX:

[particularly suitable for compounds of formula I, in which

A represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein
 X represents $-NH-C(O)-NH-$ or $-NH-C(S)-NH-$,
 Y represents phenylene or 4-8C-cycloalkylene,
 m is an integer from 0 to 4,
 n is an integer from 1 to 4];

Exemplarily shown is X = $-NH-C(S)-NH-$ and Y = phenylene:



Further compounds of formula I, whose preparation is not explicitly described in the above reaction schemes can be prepared analogously to the methods described in the following examples or according to customary preparation methods known to the person skilled in the art.

All the starting compounds are known or can be prepared according to customary preparation methods known to the person skilled in the art.

The preparation of appropriate starting compounds with a phthalazinone structure is described for example in WO98/31674, WO99/31090 or EP0934933. The preparation of appropriate starting compounds with a pyridazinone structure is described for example in EP0163965.

Suitably, the conversions are carried out analogous to methods which are familiar per se to the person skilled in the art, for example, in the manner which is described in the following examples.

It is moreover known to the person skilled in the art that if there are a number of reactive centers on a starting or intermediate compound it may be necessary to block one or more reactive centers temporarily by protective groups in order to allow a reaction to proceed specifically at the desired reaction center. A detailed description for the use of a large number of proven protective groups is found, for example, in T. W. Greene, Protective groups in organic Synthesis, John Wiley & Sons, 1991.

The substances according to the invention are isolated and purified in a manner known per se, e.g. by distilling off the solvent in vacuo and recrystallizing the residue obtained from a suitable solvent or subjecting it to one of the customary purification methods, such as column chromatography on a suitable support material.

Salts are obtained by dissolving the free compound in a suitable solvent (for example a ketone like acetone, methylethylketone, or methylisobutylketone, an ether, like diethyl ether, tetrahydrofuran or dioxane, a chlorinated hydrocarbon, such as methylene chloride or chloroform, or a low molecular weight aliphatic alcohol, such as ethanol, isopropanol) which contains the desired acid or base, or to which the desired acid or base is then added. The salts are obtained by filtering, reprecipitating, precipitating with a non-solvent for the addition salt or by evaporating the solvent. Salts obtained can be converted by basification or by acidifying into the free compounds which, in turn, can be converted into salts. In this manner, pharmacologically non-tolerable salts can be converted into pharmacologically tolerable salts.

The following examples illustrate the invention in greater detail, without restricting it. The compounds, which are mentioned in the examples as well as their salts are preferred compounds of the invention.

ExamplesFinal products

1. (cis)-2-(4-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}-amino]-1-butyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one fumarate

Prepared from compound 1a and compound A according to REACTION 2. M. p. 85-88°C

2. (cis)-2-(6-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}-amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1b and compound A according to REACTION 2. M. p. 110-112°C

3. (cis)-2-(8-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}-amino]-1-octyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1c and compound A according to REACTION 2. M. p. 111-113°C

4. (cis)-2-(6-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}-amino]-1-hexyl)-4-(3,4-diethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1d and compound A according to REACTION 2. M. p. 109-111°C

5. (cis)-2-(4-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}amino]-1-ethyl)phenyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1j and compound A according to REACTION 2. M. p. 121-125°C

6. (cis)-2-(6-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}-amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1e and compound A according to REACTION 2. M. p. 119-122°C

7. (cis)-2-(6-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-2H-phthalazin-1-one hemifumarate

Prepared from compound 1k and compound A according to REACTION 2. M. p. 107-110°C

8. (cis)-2-(2-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}amino]-1-ethyl)-4-(2,3-dihydro-7-methoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-4a,5,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 1i and compound A according to REACTION 2. M. p. 176-177°C

9. (cis)-2-(4-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}amino]-1-butyl)-4-(2,3-dihydro-7-methoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-4a,5,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1h and compound A according to REACTION 2. M. p. 120-123°C

10. (cis)-2-(6-[{2-(4-amino-3,5-dichlorophenyl)-2-hydroxyethyl}amino]-1-hexyl)-6-(3-methoxy-4-difluoromethoxyphenyl)-2H-pyridazin-3-one hemifumarate

Prepared from compound 1l and compound A according to REACTION 2. M. p. 80-85°C

11. (cis)-2-(4-[{2-(4-amino-3-chloro-5-cyanophenyl)-2-hydroxyethyl}amino]-1-butyl)-4-(3,4-dimethoxyphenyl)-4a,5,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 3b and compound D according to REACTION 5. M. p. 115-118°C.

12. (cis)-2-(5-[{2-(4-amino-3-chloro-5-cyanophenyl)-2-hydroxyethyl}amino]-1-pentyl)-4-(3,4-dimethoxyphenyl)-4a,5,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 3c and compound D according to REACTION 5. M. p. 121-123°C

13. (cis)-2-(6-[{2-(4-amino-3-chloro-5-cyanophenyl)-2-hydroxyethyl}amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 3a and compound D according to REACTION 5. M. p. 168-170°C

14. (cis)-4-(3,4-dimethoxyphenyl)-2-(6-[{2-(4-hydroxy-3-hydroxymethylphenyl)-2-hydroxyethyl}amino]-1-hexyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1b and compound C according to REACTION 2. M. p. 99-103°C

15. (cis)-2-(8-[{2-(4-hydroxy-3-hydroxymethylphenyl)-2-hydroxyethyl}amino]-1-octyl)-4-(3,4-dimethoxyphenyl)-5,6,7,8,9,10-hexahydro-1-phthalazinone 3/4fumarate

Prepared from compound 1c and compound C according to REACTION 2. M. p. 66-70°C

16. (cis)-2-(6-[{2-(4-hydroxy-3-hydroxymethylphenyl)-2-hydroxyethyl}amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 1e and compound C according to REACTION 2. M. p. 96-99°C

17. (cis)-2-(6-[{2-(4-amino-3-chloro-5-trifluoromethylphenyl)-2-hydroxyethyl}amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one 3/4fumarate

Prepared from compound 1b and compound B according to REACTION 2. M. p. 102-105°C

18. (cis)-2-(6-[{2-(4-amino-3-cyanophenyl)-2-hydroxyethyl}amino]-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one hemifumarate

Prepared from compound 4 and compound E according to REACTIONS 7, 8 and 9. M. p. 101-103°C

19. (cis)-4-(3,4-dimethoxyphenyl)-2-[6-((2-phenyl-2-hydroxyethyl)amino]-1-hexyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one 3/4fumarate

Prepared from compound 1b and 2-amino-1-phenylethanol according to REACTION 2.

M. p. 110-115°C.

20. 2-Amino-3-chloro-5-[2-(4-[4-[(4aS,8aR)-4-(3,4-dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidin-1-yl]-4-oxo-butylamino)-1-hydroxy-ethyl]-benzonitrile fumarate

Prepared from starting compound 5a and starting compound D according to REACTION 5.

M. p. 212-216°C.

21. (4aS,8aR)-2-(4-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-butoxy)-phenyl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 6a and compound A according to REACTION 2. M. p. 126-127°C.

22. (4aS,8aR)-2-(4-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-butoxy)-benzyl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 7a and compound A according to REACTION 2. M. p. 119-120°C.

23. (4aS,8aR)-2-(4-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-ethoxy)-phenyl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one oxalate

Prepared from compound 8a and compound A according to REACTION 2. M. p. 136-138°C.

24. (4aS,8aR)-2-[1-(2-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-ethanesulfonyl)-ethyl]-piperidin-4-yl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 9a and compound A according to REACTION 10. M. p. 137-138°C.

25. 4-[(4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidine-1-carbothioic acid {4-[2-(4-amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-butyl}-amide

Prepared from compound 10a and compound O according to REACTION 5. M. p. 139-144°C.

26. (4aS,8aR)-2-(1-{2-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-ethanoyl}-piperidin-4-yl)-4-(3,4-diethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 11a and compound A according to REACTION 2. M. p. 172-173°C.

27. (4aS,8aR)-2-(1-[2-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-ethanesulfonyl]-piperidin-4-yl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 12a and compound A according to REACTION 2. M. p. 169-171°C.

28. (4aS,8aR)-2-(1-[(S)-2-[2-(3,4-Diamino-5-chloro-phenyl)-2-hydroxy-ethylamino]-propanoyl]-piperidin-4-yl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 13a and compound O according to REACTION 5. M. p. 130-131°C.

29. (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-(1-[2-hydroxy-2-(4-hydroxy-3-hydroxymethyl-phenyl)-ethylamino]-ethanesulfonyl)-piperidin-4-yl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 12a and compound C according to REACTION 10. Crystallised as the free base from ethyl acetate. M. p. 97-99°C.

30. (4aS,8aR)-2-[1-(3-[2-(4-Amino-3,5-dichloro-phenyl)-2-hydroxy-ethylamino]-ethylsulfonyl)-propanoyl]-piperidin-4-yl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one hydrochloride

Prepared from compound O and compound 14a according to REACTION 5. M. p. 118-120°C.

31. (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-[4-(4-[2-hydroxy-2-(4-hydroxy-3-hydroxymethyl-phenyl)-ethylamino]-butoxy)-phenyl]-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one fumarate

Prepared from compound 6a and compound C according to REACTION 2. M. p. 126-127°C.

32. 2-Amino-3-chloro-5-[2-[3-[4-[(4aS,8aR)-4-(3,4-dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidin-1-yl]-3-oxo-propane-1-sulfonyl]-ethylamino]-1-hydroxy-ethyl]-benzonitrile

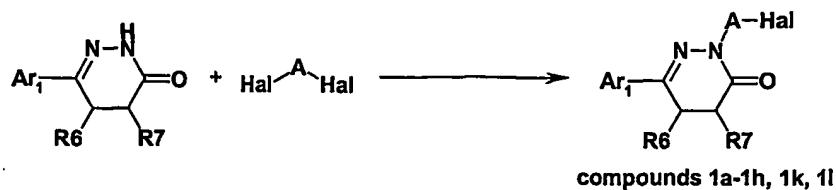
Prepared from compound 15a and compound D according to REACTION 5. Crystallised from diethyl ether as the free base. M. p. 102-103°C.

33. 2-Amino-3-chloro-5-[2-((S)-2-{4-[(4aS,8aR)-4-(3,4-dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidin-1-yl}-1-methyl-2-oxo-ethylamino)-1-hydroxy-ethyl]-benzonitrile hydrochloride

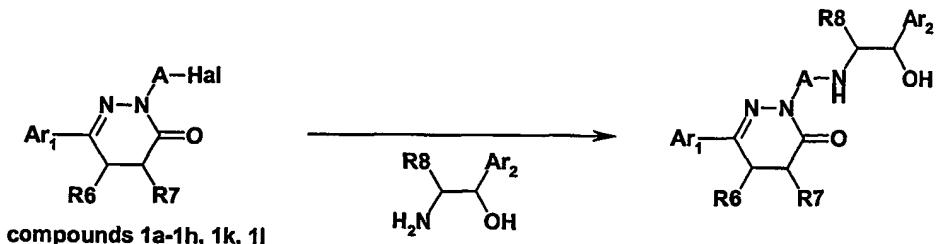
Prepared from compound 13a and compound D according to REACTION 5. Crystallised as the hydrochloride. M. p. 160-161°C.

34. 2-Amino-3-chloro-5-[2-(6-{4-[(4aS,8aR)-4-(3,4-dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidin-1-yl}-hexylamino)-1-hydroxy-ethyl]-benzonitrile fumarate

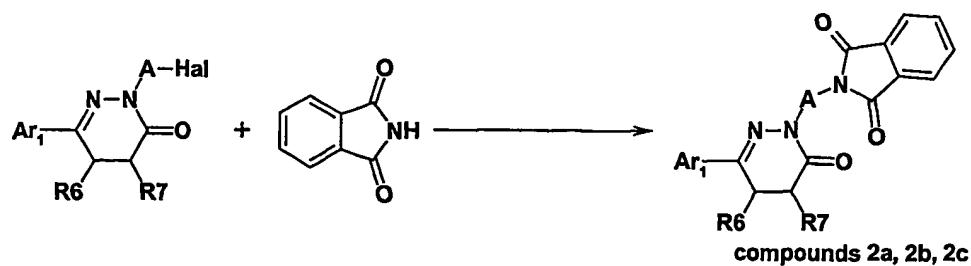
Prepared from compound 16a and compound D according to REACTION 5. M. p. 142-143°C.

GENERAL SYNTHESIS**REACTION 1: 2-(ω -halogenalkyl)phthalazinones (compounds 1a-1h, 1k or 1l)**

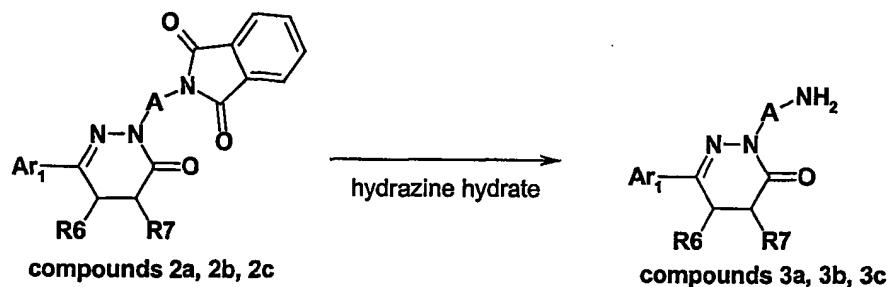
25 mmol of sodium hydride is added to a stirred solution of 20 mmol of a phthalazinone or a pyridazinone (compounds F-K) in 50 ml of dimethylformamide. After 5 min, 60 mmol of a α,ω -dihalogenalkane is added and the resulting mixture stirred for 60 min. After the addition of 200 ml of water, the mixture is extracted with diethyl ether. The ether solution is dried over magnesium sulfate and evaporated. The residue is purified by chromatography [ethyl acetate : petroleum ether (60-80°C) / 1:4].

REACTION 2: aminoethanol derivatives

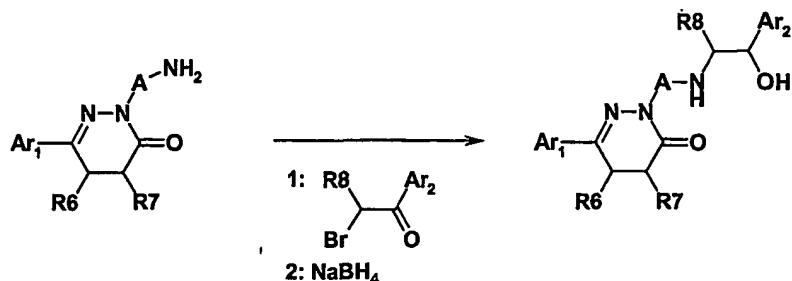
A mixture of 8 mmol of a 2-(ω -halogenalkyl)phthalazinone, 10 mmol of a primary aminoethanol derivative (compounds A, B or C) and 20 mmol of potassium carbonate in 50 ml of dimethylformamide is heated for 1.5 h at 100°C. After cooling to room temperature, a mixture of 100 ml of water and 200 ml of ethyl acetate is added. The organic layer is dried over magnesium sulfate and evaporated. The residue is purified by chromatography (ethyl acetate : triethyl amine / 20 : 1). The aminoethanol derivative is crystallized as a salt with fumaric acid from tetrahydrofuran / diethyl ether.

REACTION 3: phthalimides (compounds 2a, 2b or 2c)

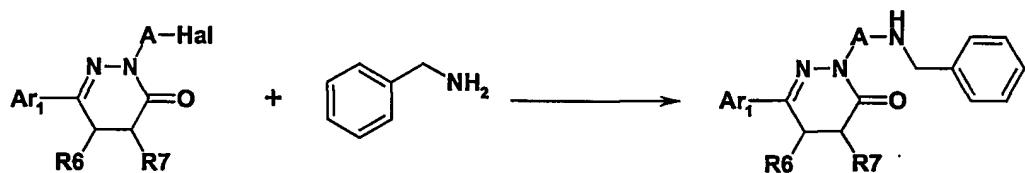
15 mmol of a 2-(ω -halogenalkyl)phthalazinone, 18 mmol of phthalimide and 20 mmol of potassium carbonate in 100 ml of dimethylformamide is heated for 5 h at 110°C. After cooling to room temperature, 300 ml of water and 300 ml of ethyl acetate is added. The organic layer is dried over magnesium sulfate and concentrated under reduced pressure. The compound is crystallized from a suitable solvent.

REACTION 4: 2-(ω -aminoalkyl)phthalazinone (compounds 3a, 3b or 3c)

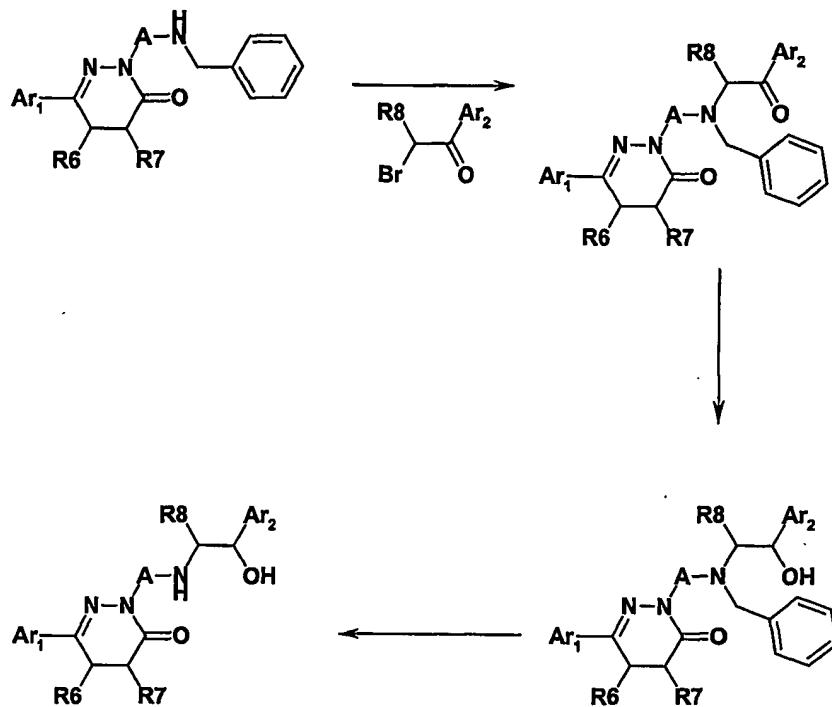
A solution of 20 mmol of a phthalimide (compounds 2a, 2b or 2c) and 25 mmol of hydrazine hydrate in 200 ml of ethanol is refluxed for 5 h and subsequently evaporated. The residue is dissolved in ethyl acetate (200 ml) and this solution is washed with aqueous sodium carbonate. After drying over magnesium sulfate, the solvent is evaporated and the residue crystallized from a suitable solvent.

REACTION 5: aminoethanol derivatives

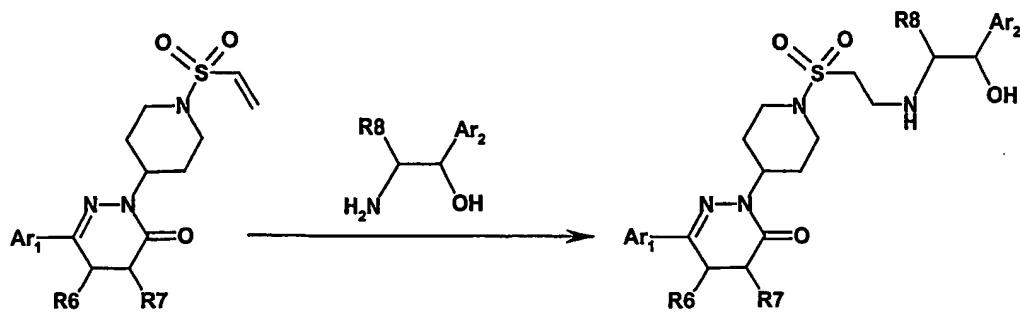
A solution of 6 mmol of a 2-bromoacetophenone derivative (for example: compounds D or E), 5 mmol of an amine (for example: compounds 3a, 3b or 3c) and 10 mmol of diisopropylethylamine in a mixture of 50 ml of tetrahydrofuran and 50 ml of methanol is stirred at room temperature. After 3 h 10 mmol of sodium borohydride is added and the resulting mixture is stirred for another hour and then evaporated. The residue is dissolved in water (100ml) and this mixture is extracted twice with 200ml of ethyl acetate. The organic solution is dried over magnesium sulfate and evaporated. The residue is purified by chromatography (ethyl acetate : methanol : triethyl amine / 10 : 2 : 0.5). The compound is crystallized as a fumarate salt from tetrahydrofuran / diethyl ether.

REACTION 6: Benzylamines

A mixture of 25 mmol of a 2-(ω -halogenalkyl)phthalazinone and 200 mmol of benzylamine is heated at 120°C for 2 h. After cooling to about 60°C, 200 ml of 2 N hydrobromic acid is added and the resulting mixture is, after cooling to room temperature, extracted twice with 200 ml of dichloromethane. The organic solution is dried over magnesium sulfate and evaporated. The residue is washed with diethyl ether and dried.

REACTIONS 7/8/9

A solution of 10 mmol of a benzylamine (compounds prepared according to REACTION 6), 10 mmol of a 2-bromoacetophenone derivative and 20 mmol of diisopropylethylamine in a mixture of 50 ml of tetrahydrofuran and 50 ml of ethanol is stirred for 18 h at room temperature after which 10 mmol of sodium borohydride is added. This mixture is stirred for another 30 min and subsequently evaporated. The residue is dissolved in 200 ml of diethyl ether and this solution is washed with aqueous sodium carbonate. The organic solution is dried over magnesium sulfate and evaporated. The residue is dissolved in ethanol and, after the addition of 0.5 g of 5% Pd/C, treated with a flow of hydrogen. After about 60 min debenzylation is completed (tlc analysis). The catalyst is filtered off, the solvent is evaporated and the residue purified by chromatography on silica (ethyl acetate : triethyl amine / 20 : 1). The aminoethanol derivative is crystallized as a fumarate salt from tetrahydrofuran / diethyl ether.

REACTION 10

A mixture of 5 mmol of an ethenesulfonyl derivative (compounds 9a or 12a) and 5 mmol of an amino-ethanol derivative (compounds A, B or C) in 50 ml of tetrahydrofuran is stirred for 18 h at RT, after which the solvent is evaporated and the residue purified by chromatography (elution with ethyl acetate containing 5% of triethylamine).

Starting Compounds and Intermediates

1a: (cis)-2-(4-bromo-1-butyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one

Prepared from compound F and 1,4-dibromobutane according to REACTION 1. M. p. oil

1H-NMR(CDCl₃): 1.28-2.00 (m, 11H, 7xcyclohexane-H, C-(CH₂)₂-C); 2.50-2.61 (m, 1H, cyclohexane-H); 2.63-2.73 (m, 1H, cyclohexane-H); 3.00-3.15 (m, 1H, cyclohexane-H); 3.49 (t, J=6.4Hz, 2H, CH₂-Br); 3.72-4.12 (m, 8H, 2xO-CH₃, N-CH₂); 6.87 (d, J=8.4Hz, 1H, Ar-H); 7.23 (dd, J=2.1, 8.5Hz, 1H, Ar-H); 7.47 (d, J=2.0Hz, 1H, Ar-H)

1b: (cis)-2-(6-bromo-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one

Prepared from compound F and 1,6-dibromohexane according to REACTION 1. Crystallized from diethyl ether. M. p. 74-75°C

1c: (cis)-2-(8-bromo-1-octyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one

Prepared from compound F and 1,8-dibromoocetane according to REACTION 1. Crystallized from hexane. M. p. 58-60°C

1d: (cis)-2-(6-bromo-1-hexyl)-4-(3,4-diethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one

Prepared from compound H and 1,6-dibromohexane according to REACTION 1. M. p. 47-51°C

1e: (cis)-2-(6-bromo-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound G and 1,6-dibromohexane according to REACTION 1. Crystallized from methanol. M. p. 55-56°C

1H-NMR(CDCl₃): 1.37-2.26 (m, 11H, 3xcyclohexene-H, C-(CH₂)₄-C); 2.67-2.81 (m, 1H, cyclohexene-H); 2.97-3.07 (m, 1H, cyclohexane-H); 3.28-3.42 (m, 3H, CH₂-Br, cyclohexene-H); 3.68-4.06 (m, 8H,

2xO-CH₃, N-CH₂); 6.88 (d, J=8.5Hz, 1H, Ar-H); 7.26 (dd, J=2.1, 8.5Hz, 1H, Ar-H); 7.48 (d, J=2.0Hz, 1H, Ar-H)

1f: (cis)-2-(4-bromo-1-butyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound G and 1,4-dibromobutane according to REACTION 1. Crystallized from methanol. M. p. 102-103°C.

1g: (cis)-2-(5-bromo-1-pentyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound G and 1,5-dibromopentane according to REACTION 1. M. p. oil.

1H-NMR(CDCl₃): 1.38-2.28 (m, 11H, 3xcyclohexene-H, C-(CH₂)₃-C); 2.67-2.81 (m, 1H, cyclohexene-H); 2.97-3.07 (m, 1H, cyclohexene-H); 3.28-3.50 (m, 3H, CH₂-Br, cyclohexene-H); 3.68-4.08 (m, 9H, 2xO-CH₃, N-CH₂, cyclohexene-H); 5.60-5.87 (m, 2H, CH=CH); 6.88 (d, J=8.5Hz, 1H, Ar-H); 7.26 (dd, J=2.0, 8.5Hz, H, Ar-H); 7.48 (d, J=2.0Hz, 1H, Ar-H).

1h: (cis)-2-(4-bromo-1-butyl)-4-(2,3-dihydro-7-methoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound I and 1,4-dibromobutane according to REACTION 1. Crystallized from methanol. M. p. 86-88°C.

1i: (cis)-2-bromoethyl-4-(2,3-dihydro-7-methoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A solution 1.92 g of Br₂ in CH₂Cl₂ is added to a solution of 3.1 g of triphenylphosphine in CH₂Cl₂ at 0°C, followed by the addition of a solution of 4.6 g of compound L in CH₂Cl₂. The resulting solution is stirred for 2 h at room temperature and subsequently washed with diluted hydrochloric acid (2x) and aqueous sodium carbonate. Crystallization from methanol (2x). M. p. 143-145°C

1j: (cis)-2-[4-(2-bromoethyl)phenyl]-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from M as described for intermediate 1i. Crystallized from methanol. M. p. 126-127°C.

1k: 2-(6-bromo-1-hexyl)-4-(3,4-dimethoxyphenyl)-2H-phthalazin-1-one

Prepared from compound J and 1,6-dibromohexane according to REACTION 1. Crystallized from a mixture of ethyl acetate and hexane. M. p. 83-84°C

1l: 2-(6-bromo-1-hexyl)-4-(3-methoxy-4-difluoromethoxyphenyl)-2H-pyridazin-1-one

Prepared from compound K and 1,6-dibromohexane according to REACTION 1. Crystallized from diethyl ether. M. p. 69-70°C.

2a: (cis)-N-[6-{4-(3,4-dimethoxyphenyl)-1-oxo-4a,5,8,8a-tetrahydro-2H-phthalazin-2-yl}hexyl]-phthalimide

Prepared from compound 1e according to REACTION 3. Crystallized from methanol. M. p. 104-105°C

2b: (cis)-N-[4-{4-(3,4-dimethoxyphenyl)-1-oxo-4a,5,8,8a-tetrahydro-2H-phthalazin-2-yl}butyl]-phthalimide

Prepared from compound 1f according to REACTION 3. Crystallized from methanol. M. p. 139-140°C.

2c: (cis)-N-[5-{4-(3,4-dimethoxyphenyl)-1-oxo-4a,5,8,8a-tetrahydro-2H-phthalazin-2-yl}pentyl]-phthalimide

Prepared from compound 1g according to REACTION 3. Crystallized from diethyl ether. M. p. 126-129°C.

3a: (cis)-2-(6-amino-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 2a according to REACTION 4. Crystallized from tetrahydrofuran / diethyl ether as a fumarate salt. M. p. 141-143°C.

3b: (cis)-2-(4-amino-1-butyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 2b according to REACTION 4. Crystallized from diethyl ether. M. p. 108-109°C.

3c: (cis)-2-(5-amino-1-pentyl)-4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 2c according to REACTION 4. Crystallized from diethyl ether. M. p. 88-89

4a: (cis)-2-(6-benzylamino-1-hexyl)-4-(3,4-dimethoxyphenyl)-4a,5,6,7,8,8a-hexahydro-2H-phthalazin-1-one HBr

Prepared from compound 1b according to REACTION 6. M. p. 117-119°C

5a: (4aS,8aR)-2-[1-(4-Amino-butanoyl)-piperidin-4-yl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one trifluoroacetate

15 mmol of compound 5b is dissolved in 20 ml of trifluoroacetic acid and the solution is left at RT for 30 min after which it is evaporated. The residue solidifies on treating with diethyl ether.

M. p. 175-178°C.

5b: (4-{4-[(4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidin-1-yl}-4-oxo-butyl)-carbamic acid tert butyl ester

To a solution of 20 mmol of compound 5c and 25 mmol of (4-Oxo-pentyl)-carbamic acid tert-butyl ester in 100 ml of dichloromethane, 25 mmol of (3-Dimethylamino-propyl)-ethyl-carbodiimide hydrochloride is added and the resulting mixture is stirred at RT. After 18 h the solution is washed successively with diluted hydrochloric acid and aqueous sodium carbonate and then dried over magnesium sulfate. After evaporating the solvent, the compound is crystallised from hexane. M. p. 79-83°C.

5c: (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-piperidin-4-yl-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A solution of 50 mmol of compound 5d in 150 ml of dichloromethane is washed with a saturated solution of sodium bicarbonate, dried over magnesium sulfate and evaporated. The residue is washed with diethyl ether and dried. M. p. 260°C (with decomposition)

5d: (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-piperidin-4-yl-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one hydrochloride

A solution of 50 mmol of the salt of (S)-(-)- α -methylbenzylamine and (cis)-2-(3,4-dimethoxybenzoyl)-1,2,3,6-tetrahydrobenzoic acid (compound N), 55 mmol of piperidin-4-yl-hydrazine dihydrochloride and 100 mmol of triethylamine in 150 ml of 1-propanol is refluxed for 18 h. After cooling to RT, the precipitate is filtered off and dried. M. p. 285-288°C

5e: Piperidin-4-yl-hydrazine dihydrochloride

A mixture of 0.1 mole of 4-(N'-tert-Butoxycarbonyl-hydrazino)-piperidine-1-carboxylic acid tert-butyl ester (compound 5f) and 150 ml of concentrated hydrochloric acid is heated at 90°C for 60 min after which the clear solution is evaporated. The residue is washed with tetrahydrofuran, filtered off and dried under vacuum. M. p. 256-259°C

5f: 4-(N'-tert-Butoxycarbonyl-hydrazino)-piperidine-1-carboxylic acid tert-butyl ester

150 ml of a solution of borohydride in tetrahydrofuran (1.0 mol/l) is slowly added to a solution of 0.12 mole of 4-(tert-Butoxycarbonyl-hydrazono)-piperidine-1-carboxylic acid tert-butyl ester (compound 5g) in 100 ml of dry tetrahydrofuran. After complete addition, the mixture is stirred for another 30 min after which a 100 ml of water is added to destroy the excess of borohydride. Subsequently the tetrahydrofuran is evaporated and the resulting aqueous solution extracted with diethyl ether. After drying the solvent over magnesium sulfate, the ether is evaporated. M. p. 112-115°C

5g: 4-(tert-Butoxycarbonyl-hydrazono)-piperidine-1-carboxylic acid tert-butyl ester

A mixture of 0.15 mole of 4-oxo-piperidine-1-carboxylic acid tert-butyl ester (commercially available) and 0.15 mole of tert-butylcarbazate in 250 ml of hexane is stirred for 18 h at RT. The precipitate is filtered off and dried under vacuum. M. p. 172-174°C

6a: (4aS,8aR)-2-[4-(4-Bromo-butoxy)-phenyl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A mixture of 10 mmol of compound 6b, 50 mmol of 1,4-dibromobutane and 40 mmol of potassium carbonate in 50 ml of dimethylformamide is stirred for 18 h at RT, after which the mixture is poured into 200 ml of water. After extraction with diethyl ether, the compound is purified by chromatography (ethyl acetate:hexane/1;5). Crystallised from hexane at -20 °C. M. p. 38-43°C.

6b: (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-(4-hydroxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A mixture of 20 mmol of compound 6c, 50 ml of ethanethiol and 20 ml of borontrifluoride etherate is left at RT for 40 h. After the addition of 200 ml of a saturated solution of sodium bicarbonate, the mixture is extracted with ethyl acetate. The organic solution is dried over magnesium sulfate and evaporated. The compound is crystallised from diethyl ether. M. p. 111-112°C.

6c: (4aS,8aR)-2-(4-Benzylxy-phenyl)-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A solution of 50 mmol of compound 6d and 50 mmol of 4-benzylxyphenylhydrazine hydrochloride in 100 ml of pyridine is refluxed for 8 h and subsequently evaporated. The residue is dissolved in ethyl acetate and the solution is washed successively with diluted hydrochloric acid and aqueous sodium carbonate and then dried over magnesium sulfate. After evaporating the solvent, the compound is crystallised from methanol. M. p. 104-105°C.

6d: (1R,6S)-6-[1-(3,4-Dimethoxy-phenyl)-methanoyl]-1,2,3,6-tetrahydrobenzoic acid

A solution of 50 mmol of compound 6e in 150 ml of dichloromethane is washed successively with 1 N sulfuric acid and water (twice). The solution is dried over magnesium sulfate and evaporated. The residue is crystallised from diethyl ether. M. p. 110-112°C.

6e: (1R,2S)-2-(3,4-Dimethoxy-phenyl)-methanoyl)-1,2,3,6-tetrahydrobenzoic acid (-)- α -methylbenzylamine salt

A mixture of 10 mmol of compound N and 5 mmol of L- α -methylbenzylamine in 100 ml of ethyl acetate is stirred for 18 h after which the precipitate is filtered off and dried.

7a: (4aS,8aR)-2-[4-(4-Bromo-butoxy)-benzyl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 7b and 1,4-dibromobutane as described for compound 6a. M. p. 42-46°C.

7b: (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-(4-hydroxy-benzyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from 4-hydroxybenzylhydrazine hydrochloride and compound 6d as described for compound 6c. Crystallised from diethyl ether. M. p. 189-190°C.

8a: (4aS,8aR)-2-[4-(2-Bromo-ethoxy)-phenyl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 6b and 1,2-dibromoethane as described for compound 6a. M. p. 40-45°C.

9a: (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-[1-(2-ethenesulfonyl-ethyl)-piperidin-4-yl]-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A solution of 10 mmol of compound 5c, 10 mmol of 1-Bromo-2-(2-bromo-ethanesulfonyl)-ethane and 30 mmol of potassium carbonate in 30 ml of dimethylformamide is stirred at RT. After 18 h, the mixture is poured into water (150 ml) and subsequently extracted with ethyl acetate. Crystallisation from diethyl ether. M. p. 165-166°C

10a: 4-[(4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1H-phthalazin-2-yl]-piperidine-1-carbothioic acid (4-amino-butyl)-amide trifluoroacetate

Prepared from compound 10b as described for compound 5a. M. p. 125-128°C.

10b: {4-[(1-(4-[(4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1-phthalazin-2-yl]-piperidin-1-yl)-methanethioyl]-amino]-butyl}-carbamic acid tert butyl ester

A solution of 10 mmol of (4-Isothiocyanato-butyl)-carbamic acid tert butyl ester and 10 mmol of compound 5c in 100 ml of dichloromethane is stirred for 18 h at RT, successively washed with diluted hydrochloric acid and aqueous sodium carbonate and subsequently dried over magnesium sulfate and evaporated. Crystallised from a mixture of ethyl acetate and hexane. M. p. 118-121°C.

11a: (4aS,8aR)-2-[1-(2-Chloro-ethanoyl)-piperidin-4-yl]-4-(3,4-diethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A solution of 12 mmol of chloroacetylchloride in 20 ml of dichloromethane is added at 0°C to a mixture of 12 mmol of compound 11b and 40 mmol of triethylamine in 50 ml of dichloromethane. The resulting mixture is stirred for 60 min at RT and subsequently washed with diluted hydrochloric acid and aqueous sodium carbonate. After drying the solution over magnesium sulfate, the solvent is evaporated and the residue purified by chromatography (ethyl acetate:petroleum ether (60-80°C)/2:1). M. p. 135-136°C.

11b: (4aS,8aR)-4-(3,4-Diethoxy-phenyl)-2-piperidin-4-yl-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one hydrochloride

Prepared from the salt of (S)-(-)- α -methylbenzylamine, (cis)-2-(3,4-diethoxybenzoyl)-1,2,3,6-tetrahydrobenzoic acid (compound P) and piperidin-4-yl-hydrazine dihydrochloride in 2-propanol as described for compound 5d. M. p. 248-250°C

12a: (4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-2-(1-ethenesulfonyl-piperidin-4-yl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

A solution of 20 mmol of 2-chloroethylsulfonylchloride in 20 ml of dichloromethane is added to a solution of 15 mmol of compound 5c and 25 ml of diisopropylethylamine in 50 ml of dichloromethane. The resulting mixture is stirred at RT for 18 h, after which it is washed first with diluted hydrochloric acid and then with a saturated solution of sodium bicarbonate. After drying over magnesium sulfate and evaporating the solvent, the compound is crystallised from diethyl ether. M. p. 89-90°C.

13a: (4aS,8aR)-2-[1-((S)-2-Amino-propanoyl)-piperidin-4-yl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared from compound 13b as described for compound 5a. After evaporating the trifluoroacetic acid, the residue is partitioned between aqueous sodium carbonate and dichloromethane. The organic solution is dried over magnesium sulfate and evaporated. The residue is crystallised from diethyl ether. M. p. 101-103°C.

13b: 1-tert-Butyl-3-(*S*)-2-{4-[(4a*S*,8a*R*)-4-(3,4-dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1*H*-phthalazin-2-yl]-piperidin-1-yl}-1-methyl-2-oxo-ethyl)-urea

Prepared from compound 5c and (R)-2-(3-tert-Butyl-ureido)-propionic acid (N-Boc-L-alanine) as described for compound 5b. M. p. 153-154°C.

14a. (4a*S*,8a*R*)-2-{1-[3-(2-Amino-ethylsulfanyl)-propanoyl]-piperidin-4-yl}-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2*H*-phthalazin-1-one

Prepared from compound 14b as described for compound 5c. After evaporating the trifluoroacetic acid, the residue is partitioned between a bicarbonate solution and dichloromethane. The dichloromethane solution is dried over magnesium sulfate and evaporated. Crystallised from ethyl acetate / hexane. M. p. 137-139°C.

14b. [2-{3-[(4a*S*,8a*R*)-4-(3,4-Dimethoxy-phenyl)-1-oxo-4a,5,8,8a-tetrahydro-1*H*-phthalazin-2-yl]-piperidin-1-yl}-3-oxo-propylsulfanyl]-ethyl]-carbamic acid tert butyl ester

Prepared from 12 mmol of 3-(2-tert-Butoxycarbonylamino-ethylsulfanyl)-propionic acid and 10 mmol of compound 5c as described for compound 5b. M. p. 132-134°C.

15a. (4a*S*,8a*R*)-2-{1-[3-(2-Amino-ethanesulfonyl)-propanoyl]-piperidin-4-yl}-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2*H*-phthalazin-1-one hydrochloride

20 mmol of 3-chloroperbenzoic acid is added to a solution of 10 mmol of compound 14a in 50 ml of dichloromethane. The resulting mixture is stirred for 30 min at RT and subsequently washed with aqueous sodium carbonate. After drying over magnesium sulfate, a solution of hydrochloric acid in diethyl ether is added to the dichloromethane solution. The precipitate is filtered off and dried. M. p. 149-150°C.

16a. (4a*S*,8a*R*)-2-[1-(6-Amino-hexyl)-piperidin-4-yl]-4-(3,4-dimethoxy-phenyl)-4a,5,8,8a-tetrahydro-2*H*-phthalazin-1-one hydrochloride

Prepared from compound 16b as described in REACTION 4. Crystallised from ethyl acetate as the hydrochloride. M. p. 176-177°C.

16b. 2-(6-[(4aS,8aR)-4-(3,4-Dimethoxy-phenyl)-1-oxo-4a,5,8a-tetrahydro-1H-phthalazin-2-yl]-piperidin-1-yl)-hexyl)-isoindole-1,3-dione

A mixture of 6 mmol of 2-(6-Bromo-hexyl)-isoindole-1,3-dione, 6 mmol of compound 5c and 3 g of potassium carbonate in 50 ml of dimethylformamide is stirred for 18 h at RT. After evaporating the solvent, the residue is partitioned between water and ethyl acetate. The organic layer is dried over magnesium sulfate and evaporated. The residue is purified by chromatography (ethyl acetate:hexane/4:1). M. p. 39-44°C.

A: 2-amino-1-(4-amino-3,5-dichlorophenyl)ethanol

Prepared as described in J. Keck, G. Kruger, K. Noll and H. Machleidt; Arzneim.-Forsch. (Drug Res.) 22, 861-869 (1972)

B: 2-amino-1-(4-amino-3-chloro-5-trifluoromethylphenyl)ethanol

Prepared as described in G. Krüger, J. Keck, K. Noll and H. Pieper; Arzneim.-Forsch./Drug Res. 34, 1612-1624 (1984).

C: 2-amino-1-(4-hydroxy-3-hydroxymethylphenyl)ethanol

Prepared as described in D.T. Cullin, D. Hartley, L.H.C. Lunts, J.C. Press, A.C. Ritchie and P. Toon; J. Med. Chem., 13, 674-680 (1970).

D: 4'-amino-3'-chloro-5'-cyano-2-bromoacetophenone

Prepared as described in G. Krüger, J. Keck, K. Noll and H. Pieper; Arzneim.-Forsch./Drug Res. 34, 1612-1624 (1984).

E: 4'-amino-3'-cyano-2-bromoacetophenone

Prepared as described in G. Krüger, J. Keck, K. Noll and H. Pieper; Arzneim.-Forsch./Drug Res. 34, 1612-1624 (1984).

F: 4-(3,4-dimethoxyphenyl)-4a,5,6,7,8a-hexahydro-2H-phthalazin-1-one

Prepared as described in example 1 of WO98/31674.

G: 4-(3,4-dimethoxyphenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared as described in example 3 of WO98/31674.

H: 4-(3,4-diethoxyphenyl)-4a,5,6,7,8,8a-hexahydro-2H-phthalazin-1-one

Prepared as described in example 7 of WO98/31674.

I: (cis)-4-(2,3-dihydro-7-methoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared as described in example 4 of WO99/31090.

J: 4-(3,4-dimethoxyphenyl)-2H-phthalazin-1-one

Prepared as described in example B of EP0934933.

K: 4-(4-Difluoromethoxy-3-methoxyphenyl)-2H-pyridazin-1-one

Prepared as described in example 4 of EP0163965.

L: (cis)-4-(2,3-dihydro-7-methoxybenzofuran-2-spiro-1'-cyclopentan-4-yl)-2-hydroxyethyl-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared as described in example 17 of WO99/31090

M: (cis)-4-(3,4-dimethoxyphenyl)-2-(4-(2-hydroxyethyl)phenyl)-4a,5,8,8a-tetrahydro-2H-phthalazin-1-one

Prepared as described in example 70 of WO98/31674.

N. (cis)-2-(3,4-Dimethoxybenzoyl)-1,2,3,6-tetrahydrobenzoic acid

Prepared as described in WO98/31674.

O. 1-(4-Amino-3,5-dichloro-phenyl)-2-bromo-ethanone

Prepared as described by J. Keck, G. Krüger, K. Noll and H. Machleidt; Arzneim.-Forsch. (Drug Res.) 22, 861-869 (1972).

P. (cis)-2-(3,4-diethoxybenzoyl)-1,2,3,6-tetrahydrobenzoic acid

Prepared analogously as described for compound N.

Commercial applicability

β -Sympathomimetics (β 2-adrenoreceptor agonists) represent the preferred drugs for the acute relief of bronchospasm in patients with bronchial asthma (Nelson HS, β -Adrenergic bronchodilators. The New England of Medicine 333: 499-506, 1995). Although the major action of β -sympathomimetics on the airways is relaxation of airway smooth muscle, inhibitory effects on inflammatory cells (e.g. mast cells, eosinophils, neutrophils, macrophages and T-lymphocytes) in vitro have been reported (Barnes PJ, Effect of beta-agonists on inflammatory cells. Journal of Allergy and Clinical Immunology 104: S10-S17, 1999). Despite the inhibitory effects on inflammatory cells in vitro, however, β -sympathomimetics do not appear to reduce the chronic inflammation of asthma. Desensitization to the action of β -sympathomimetics is more readily seen in inflammatory cells than in airway smooth muscle cells and may account for this discrepancy. It has been suggested that an accelerated degradation of cAMP by phosphodiesterase 4 (PDE4) could be an explanation for this kind of desensitization (Giembycz MA, Phosphodiesterase 4 and tolerance to β 2-adrenoceptor agonists in asthma. TiPS 17: 331-336, 1996). Therefore, the combined action of one molecule to inhibit PDE4 activity and to activate the β 2-adrenoceptor is claimed to counteract the development of tolerance to β 2-adrenoceptor agonists in vivo. In addition, such molecules with a combined mode of action will maintain the well-known anti-inflammatory potential of PDE4 inhibitors (Torphy TJ, Phosphodiesterase isozymes – molecular targets for novel antiasthma agents. Am J Respir Crit Care Med 157: 351-370, 1998), or even make the PDE4 inhibiting component by the β 2-adrenoceptor agonistic component (more) effective under conditions where a PDE4 inhibitor requires an additional cAMP trigger for its action. One example for the latter possibility is the human eosinophil which upon stimulation by complement C5a does not respond to PDE4 inhibitors alone, but is inhibited in a highly synergistic fashion by the combined use of PDE4 inhibitors and a β 2-adrenoceptor agonist like salbutamol (Hatzelmann A, Tenor H and Schudt C, Differential effects of non-selective and selective phosphodiesterase inhibitors on human eosinophil function. Brit J Pharmacol 114: 821-831, 1995).

The compounds according to the invention have valuable pharmacological properties which make them commercially utilizable. As compounds providing selective PDE4 inhibition combined with β 2-adrenoceptor agonist activity they are suitable on the one hand as bronchial therapeutics (for the treatment of airway obstructions on account of their dilating but also on account of their respiratory rate- or respiratory drive- increasing action), but on the other hand especially for the treatment of disorders, in particular of inflammatory nature, e.g. of the airways, of the skin, of the central nervous system, of the intestine, of the eyes and of the joints, which are mediated by mediators such as histamine, PAF (platelet-activating factor), arachidonic acid metabolites such as leukotrienes and prostaglandins, cytokines, interleukins, chemokines, alpha-, beta- and gamma-interferon, tumor necrosis factor (TNF) or oxygen radicals and proteases.

On account of their PDE4-inhibiting properties, the compounds according to the invention can be employed in human and veterinary medicine and therapeutics, where they can be used, for example, for the treatment and prophylaxis of the following illnesses: acute and chronic (in particular inflammatory and allergen-induced) airway disorders of various origins (bronchitis, allergic bronchitis, bronchial asthma, emphysema, COPD); dermatoses (especially of proliferative, inflammatory and allergic type) such as, for example, psoriasis (vulgaris), toxic and allergic contact eczema, atopic eczema, seborrheic eczema, lichen simplex, sunburn, pruritus in the anogenital area, alopecia areata, hypertrophic scars, discoid lupus erythematosus, follicular and wide-area pyodermias, endogenous and exogenous acne, acne rosacea and other proliferative, inflammatory and allergic skin disorders; disorders which are based on an excessive release of TNF and leukotrienes, e.g. disorders of the arthritis type (rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis and other arthritic conditions), disorders of the immune system (AIDS, multiple sclerosis), graft-versus-host reactions, transplant rejection reactions, symptoms of shock [septic shock, endotoxin shock, gram-negative sepsis, toxic shock syndrome and ARDS (adult respiratory distress syndrome)], and generalized inflammations in the gastrointestinal area (Crohn's disease and ulcerative colitis); disorders which are based on allergic and/or chronic, faulty immunological reactions in the area of the upper airways (pharynx, nose) and the adjacent regions (paranasal sinuses, eyes), such as, for example, allergic rhinitis/sinusitis, chronic rhinitis/sinusitis, allergic conjunctivitis and nasal polyps; but also disorders of the heart which can be treated by PDE inhibitors, such as, for example, cardiac insufficiency, or disorders which can be treated on account of the tissue-relaxant action of the PDE inhibitors, such as, for example, erectile dysfunction or colics of the kidneys and the ureters in connection with kidney stones. In addition, the compounds according to the invention can be employed for the treatment of diabetes insipidus and disorders in connection with disturbances of brain metabolism, such as, for example, cerebral senility, senile dementia (Alzheimer's dementia), multiinfarct dementia or alternatively disorders of the CNS, such as, for example, depressions, arteriosclerotic dementia, Alzheimer's disease or multiple sclerosis.

A further subject of the invention is a process for the treatment of mammals, including man, which are suffering from one of the above-mentioned illnesses. The process comprises administering to the sick mammal a therapeutically efficacious and pharmacologically tolerable amount of one or more of the compounds according to the invention.

The invention further relates to the compounds according to the invention for use in the treatment of mammals, including man, which are suffering from one of the above-mentioned illnesses. The process comprises administering to the sick mammal a therapeutically efficacious and pharmacologically tolerable amount of one or more of the compounds according to the invention.

The invention further relates to the compounds according to the invention for use in the treatment and/or prophylaxis of illnesses, in particular the illnesses mentioned.

The invention likewise relates to the use of the compounds according to the invention for the production of medicaments which are employed for the treatment and/or prophylaxis of the illnesses mentioned.

Medicaments for the treatment and/or prophylaxis of the illnesses mentioned, which contain one or more of the compounds according to the invention, are furthermore a subject of the invention.

A further subject of the invention is a commercial product, consisting of a customary secondary pack, a primary pack containing the medicament (for example an inhaler or a blister pack) and, optionally, a pack insert, the medicament exhibiting simultaneously antagonistic action against cyclic nucleotide phosphodiesterases of type 4 (PDE4) and agonistic action on β 2-adrenoreceptors leading to the attenuation of the symptoms of illnesses which are connected with cyclic nucleotide phosphodiesterases of type 4 and β 2-adrenoreceptors, and the suitability of the medicament for the prophylaxis or treatment of illnesses which are connected with cyclic nucleotide phosphodiesterases of the type 4 and β 2-adrenoreceptors being indicated on the secondary pack and/or on the pack insert of the commercial product, and the medicament containing one or more compounds of the formula I according to the invention. The secondary pack, the primary pack containing the medicament and the pack insert otherwise comply with what would be regarded as standard to the person skilled in the art for medicaments of this type.

The medicaments are prepared by processes which are known *per se* and familiar to the person skilled in the art. As medicaments, the compounds according to the invention (= active compounds) are either employed as such, or preferably in combination with suitable pharmaceutical excipients, e.g. in the form of tablets, coated tablets, capsules, suppositories, patches, emulsions, suspensions, gels or solutions, the active compound content advantageously being between 0.1 and 95%.

The person skilled in the art is familiar on the basis of his/her expert knowledge with the excipients which are suitable for the desired pharmaceutical formulations. In addition to solvents, gel-forming agents, ointment bases and other active compound vehicles, it is possible to use, for example, antioxidants, dispersants, emulsifiers, preservatives, solubilizers, permeation promoters or complex formers.

For the treatment of disorders of the respiratory tract, the compounds according to the invention are preferably also administered by inhalation.

Aerosol particles in solid, liquid or mixed composition with a diameter from 0.5 to 10 µm, preferably 2 to 6 µm, are preferably used, if the compounds of the invention are administered by inhalation. The aerosol generation can be effected, for example, through pressure driven jet nebulizers or ultrasonic nebulizers, preferably however through propellant driven metered dose aerosols or through propellant free use of micronized active substances out of inhalation capsules.

Dependant on the used inhalation system the inhalation preparations contain additional to the active substance(s) further required excipients, such as propellants (for example Frigen in metered dose inhalers), surfactants, emulgators, stabilisators, preservatives, flavouring agents or fillers (for example lactose in powder inhalers).

For the purpose of inhalation a variety of devices are available with which aerosols with an optimal particle size can be generated and administered in a manner highly adapted to the patients. Beside the use of adapters (Spacer, Expander) and pear-shaped containers (for example Nebulator®, Volumatic®), as well as automatic spray burst shutter (Autohaler®) for metered dose inhalers, in particular in the field the powder inhalers there exists a variety of technical solutions (for example Diskhaler®, Rotadisk®, Turbohaler® or the inhaler described in EP 0 505321) with which an optimal administration of the active substance(s) can be achieved.

For the treatment of dermatoses, the compounds according to the invention are in particular used in the form of those medicaments which are suitable for topical application. For the production of the medicaments, the compounds according to the invention (= active compounds) are preferably mixed with suitable pharmaceutical excipients and further processed to give suitable pharmaceutical formulations. Suitable pharmaceutical formulations which may be mentioned are, for example, powders, emulsions, suspensions, sprays, oils, ointments, fatty ointments, creams, pastes, gels or solutions.

The medicaments according to the invention are prepared by methods known per se. Dosage of the active compounds takes place in the order of magnitude customary for PDE inhibitors. Thus topical application forms (such as, for example, ointments) for the treatment of dermatoses contain the active compounds in a concentration of, for example, 0.1-99%. The dose for administration by inhalation is customarily between 0.1 and 3 mg per day. The customary dose in the case of systemic therapy (p.o. or i.v.) is between 0.03 and 3 mg per kilogram per day.

Biological investigations

In the investigation of the combined action of the compounds (PDE4 inhibition and β 2-adrenoceptor agonism) at the cellular level, the activation of inflammatory cells has particular importance. As an example, the complement C5a-induced superoxide production of eosinophilic granulocytes may be mentioned, which can be measured as luminol-potentiated chemiluminescence (Hatzelmann A, Tenor H and Schudt C, Differential effects of non-selective and selective phosphodiesterase inhibitors on human eosinophil function. *Brit J Pharmacol* 114: 821-831, 1995). In this experimental setting PDE4 inhibitors by themselves are ineffective unless β 2-adrenoceptors are stimulated simultaneously.

The second messenger cyclic AMP (cAMP) is well-known for inhibiting inflammatory and immunocompetent cells. The PDE4 isoenzyme is broadly expressed in cells involved in the initiation and propagation of inflammatory diseases (H Tenor and C Schudt, in „Phosphodiesterase Inhibitors“, 21-40, „The Handbook of Immunopharmacology“, Academic Press, 1996), and its inhibition leads to an increase of the intracellular cAMP concentration and thus to the inhibition of cellular activation (JE Souness et al., *Immunopharmacology* 47: 127-162, 2000).

The antiinflammatory potential of PDE4 inhibitors *in vivo* in various animal models has been described (MM Teixeira, *TiPS* 18: 164-170, 1997). For the investigation of PDE4 inhibition on the cellular level (*in vitro*), a large variety of proinflammatory responses can be measured. Examples are the superoxide production of neutrophilic (C Schudt et al., *Arch Pharmacol* 344: 682-690, 1991) or eosinophilic (A Hatzelmann et al., *Brit J Pharmacol* 114: 821-831, 1995) granulocytes, which can be measured as luminol-enhanced chemiluminescence, or the synthesis of tumor necrosis factor- α in monocytes, macrophages or dendritic cells (Gantner et al., *Brit J Pharmacol* 121: 221-231, 1997, and *Pulmonary Pharmacol Therap* 12: 377-386, 1999). In addition, the immunomodulatory potential of PDE4 inhibitors is evident from the inhibition of T-cell responses like cytokine synthesis or proliferation (DM Essayan, *Biochem Pharmacol* 57: 965-973, 1999). Substances which inhibit the secretion of the aforementioned proinflammatory mediators are those which inhibit PDE4. The antiinflammatory action of compounds having a PDE4 inhibitory component may be enhanced by the β 2-adrenoceptor agonistic component in additive or synergistic fashion.

Method for measuring inhibition of PDE4 activity

PDE4 activity was determined as described by Thompson et al. (*Adv Cycl Nucl Res* 10: 69-92, 1979) with some modifications (Bauer and Schwabe, *Naunyn-Schmiedeberg's Arch Pharmacol* 311: 193-198, 1980). At a final assay volume of 200 μ l (96well microtiter plates) the assay mixture contained 20 mM

Tris-HCl (pH 7.4), 5 mM MgCl₂, 0.5 µM cAMP, [³H]cAMP (about 30,000 cpm/assay), the test compound and an aliquot of cytosol from human neutrophils which mainly contains PDE4 activity as described by Schudt et al. (Naunyn-Schmiedeberg's Arch Pharmacol 344: 682-690, 1991); the PDE3-specific inhibitor Motapizone (1 µM) was included to suppress PDE3 activity originating from contaminating platelets.

For (some of) the examples 1-19 stock solutions of the compounds (2 mmol/l) were prepared in DMSO and diluted 1:100 (v/v) in the Tris-HCl buffer mentioned above; appropriate dilutions were prepared in 1% (v/v) DMSO/Tris-HCl which were diluted 1:2 (v/v) in the assays to obtain the final concentrations of the inhibitors at a DMSO concentration of 0.5 % (v/v) which by itself did not affect PDE activity.

To minimize problems with the solubility of the compounds, starting from example 20 serial dilutions of the compounds were prepared directly in DMSO and further diluted 1:100 (v/v) in the assays to obtain now the final concentrations of the inhibitors at a DMSO concentration of 1 % (v/v) which by itself only slightly affected PDE4 activity.

After preincubation for 5 min at 37°C, the reaction was started by the addition of substrate (cAMP) and the assays were incubated for further 15 min at 37°C. 50 µl of 0.2 N HCl was added to stop the reaction and the assays were left on ice for about 10 min. Following incubation with 25 µg 5'-nucleotidase (Crotalus atrox snake venom) for 10 min at 37°C, the assays were loaded on QAE Sephadex A-25 (1 ml bed volume). The columns were eluted with 2 ml of 30 mM ammonium formate (pH 6.0) and the eluate was counted for radioactivity. Results were corrected for blank values (measured in the presence of denatured protein) which were below 5 % of total radioactivity. The amount of cyclic nucleotides hydrolyzed did not exceed 30 % of the original substrate concentration. The IC₅₀-values for the compounds according to the invention for the inhibition of the PDE4 activity were determined from the concentration-inhibition curves by nonlinear-regression.

The PDE4-inhibitory values determined for the compounds according to the invention [inhibitory concentration as -log IC₅₀ (mol/l)] follow from Table A below, in which the numbers of the compounds correspond to the numbers of the examples.

b) β2-Adrenoceptor agonistic activity in isolated guinea-pig trachea

Male guinea-pigs (350-450 g) were killed by a sharp blow on the head and exsanguination, after that the trachea was removed. After clearing off adhering tissue the trachea was cut into single rings which were tied together forming up to six 4- to 5-ring chains which were suspended in 10-ml organ baths containing Krebs buffer of the following composition (mM): NaCl 118, KCl 5.2, CaCl₂ 1.9, MgSO₄ 0.56, NaH₂PO₄ 0.8, NaHCO₃ 25.0, and glucose 11.1, maintained at 37°C and continuously aerated with a

mixture of 95% O₂ and 5% CO₂. The end of the chain was tied at the bottom of the tissue bath and connected to a force-displacement transducer (Type K-30, Hugo Sachs Elektronik, Germany) for the recording of isometric tension changes and then placed under 1.5 to 2 g of tension. After a 60 min equilibration period, during which time the preparations were repeatedly washed and the spontaneous contraction plateaued, each tracheal preparation was pre-treated with 3x10⁻⁷ M of RP 73401 for total inhibition of PDE4, which caused relaxation of the tissue between 30 and 40% of maximum (Harris et al., 1989, JPET 251: 199-206). After stabilization of the relaxant response to RP 73401, the test drugs were administered in a cumulative manner (factor 3), each concentration remaining in contact with the trachea until the contractile response reached a plateau, before addition of the next higher concentration. At the end of the test drug administration, which caused maximal relaxation the tissue (verified in control experiments for each test drug by additional administration of 10⁻⁶ M (-)-isoprenaline), a high concentration of the β -adrenoceptor antagonist, dl-sotalol (3-6x10⁻⁴ M), was additionally given to assess whether the relaxant response of the trachea to test compound was due to β_2 -adrenoceptor stimulation. dl-Sotalol nearly completely reversed the relaxation amplitude to the test compounds and was taken as evidence for their β_2 -adrenoceptor agonistic property. The relaxant response to test compound was expressed as percent determined between the relaxant amplitude after that reached by RP 73401 (0%) and the maximal relaxation obtained prior to dl-sotalol administration (100%). EC₅₀ values for a half-maximal relaxation of the tissue by test compound were graphically interpolated from 12 to 16 individual concentration-response curve. The -log EC₅₀ values (mol/l) thus obtained were taken as a measure for β_2 -adrenoceptor agonistic activity of the test compounds.

The -log EC₅₀ values (mol/l) for the compounds according to the invention follow from the Table A below, in which the numbers of the compounds correspond to the numbers of the examples.

c) Inhibition of C5a-stimulated chemiluminescence in human eosinophils

Eosinophils were purified essentially as described in detail elsewhere (Hatzelmann A, Tenor H and Schudt C, Differential effects of non-selective and selective phosphodiesterase inhibitors on human eosinophil functions. Brit J Pharmacol 114: 821-831, 1995). Briefly, total granulocytes were first purified from blood (anticoagulated with 0.3 % w/v sodium citrate) by dextran sedimentation, centrifugation on Ficoll Paque and hypotonic lysis of remaining red blood cells. For the further purification of the eosinophil fraction the magnetic cell separation (MACS) system from Miltenyi Biotec (Bergisch-Gladbach, Germany) was applied. Eosinophils were separated from neutrophils by negative selection using anti-CD16 microbeads in a two-step protocol using D- and BS- (formerly called B₂-) separation columns. By this method human eosinophils with a purity of > 99 % and a viability of > 97 % were obtained.

The chemiluminescence (CL) assay was essentially performed as described (Hatzelmann et al., 1995). Briefly, measurements were performed in a "CL-buffer" (140 mM NaCl, 5 mM KCl, 1 mM MgCl₂, 10 mM HEPES, pH 7.4) containing 1 mM CaCl₂, 1 mg/ml glucose, 0.05 % (w/v) BSA, 10 µM luminol and 4 µM microperoxidase (all values correspond to final concentrations in the assays) in a total volume of 0.5 ml at a cell concentration of 10⁶ cells/ml. Aliquots (0.4 ml) of the cell suspension (containing luminol and microperoxidase) were preincubated for 5 min at 37°C in the absence or presence of inhibitors (0.05 ml). Stock solutions of the inhibitors (10 mM) were prepared in DMSO and diluted 1:100 (v/v) in CL-buffer; subsequent dilutions were made in 1 % DMSO/CL-buffer to achieve the final drug concentrations in the assays at a DMSO concentration of 0.1 % (v/v) which by itself only weakly affected the CL response. As a further addition (0.01 ml) during the preincubation of the cells salbutamol (100 nM final concentration, which by itself hardly affected the CL response) was included as additional cAMP trigger. After preincubation, the assays were transferred into a "Multi-Biolumat LB 9505C" from Berthold (Wildbad, Germany) and stimulated by the addition of 0.05 ml C5a (dissolved in CL-buffer; 100 nM final concentration). CL was continuously recorded for 1 min (C5a) and the AUC (area under the curve) calculated. Inhibition of CL by the compounds under these conditions is given as $-\log IC_{50}$ (mol/l).

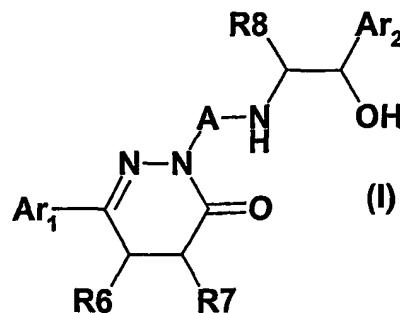
The inhibition of CL by the compounds according to the invention [as $-\log IC_{50}$ (mol/l)] follow from the Table A below, in which the numbers of the compounds correspond to the number of the examples.

Table A:

Compound	Inhibition of PDE4 activity [-logIC ₅₀ (mol/l)]	β2-Adrenoceptor agonistic affinity [-logEC ₅₀ (mol/l)]	Inhibition of C5a-stimulated chemiluminescence [-logIC ₃₀ (mol/l)]
1	6.97	7.89	5.52
2	7.33	6.86	<5
3	7.47	6.77	6.52
5	8.31		6.70
6	7.69	7.60	6.00
8	8.18		6.40
11	7.59	8.11	6.22
12	7.65	8.61	5.92
13	7.80	6.91	6.52
14	7.43	7.24	5.30
20	10.17	8.64	7.10
21	10.03		6.52
22	9.79		6.70
25	10.62		6.10
27	11.20		6.40
30	10.67		6.70
32	11.40		6.64
33	10.07		6.30

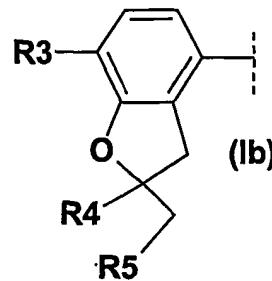
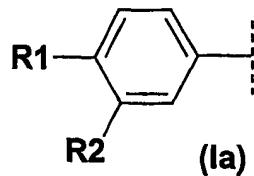
Patent claims

1. Compounds of formula I



in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

R1 is hydroxyl, halogen, 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

R2 is hydroxyl, halogen, 1-8C-alkoxy, 3-7C-cycloalkoxy, 3-7C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

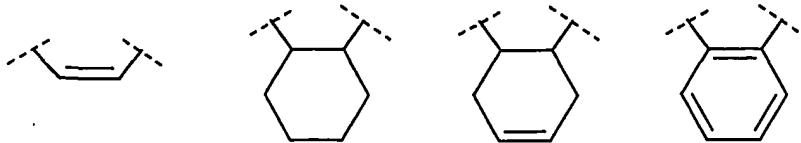
R4 is 1-4C-alkyl and

R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked 5-, 6- or 7-membered hydrocarbon ring, optionally interrupted by an oxygen or sulphur atom,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents $-C_mH_{2m}Y-X-C_nH_{2n}$ or $-Y-X-C_mH_{2m}Z-C_nH_{2n}$, wherein

X represents a bond, -O- (oxygen), -S- (sulfur), -NH-, -C(O)-, -S(O)₂-, -C(O)-NH-, -NH-C(O)-, -C(S)-NH-, -NH-C(S)-, -NH-C(O)-NH- or -NH-C(S)-NH-,

Y represents a bond, phenylene, 4-8C-cycloalkylene or azacycloalkylene,

Z represents -O-, -S-, -S(O)₂-, -NH-C(O)-, -C(O)-NH-, -NH-C(S)- or -C(S)-NH-,

m is an integer from 0 to 4,

n is an integer from 1 to 4,

R8 is hydrogen or 1-4C-alkyl,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11,

wherein

R9 is hydrogen, halogen, hydroxyl, amino, ureido [-NH-C(O)-NH₂], formylamino [-NH-C(O)H], 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,

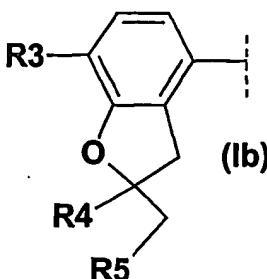
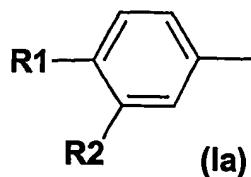
R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl or 1-4C-alkylcarbonyloxy,

R11 is hydrogen or halogen,

and the salts of these compounds.

2. Compounds of formula I according to claim 1 in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



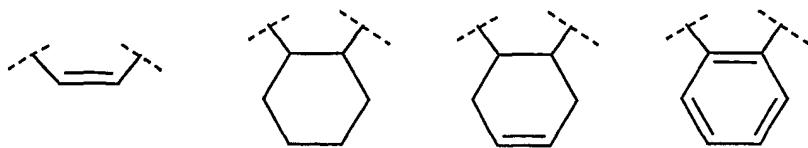
wherein

- R1 is 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is 1-4C-alkyl and
- R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane, cyclohexane, tetrahydrofuran or tetrahydropyran ring,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



and A either represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein either

- X represents a bond,
- Y represents a bond,
- m is an integer from 1 to 4, and
- n is an integer from 1 to 4,
- or
- X represents a bond, -O-, -S-, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,
- Y represents 1,4-phenylene, 1,3-phenylene, 1,4-cyclohexylene, 1,3-cyclohexylene or 1,3-cyclopentylene,
- m is an integer from 0 to 4, and

n is an integer from 1 to 4,

or

X represents a bond, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,

Y represents 4,1-piperidinylene,

m is 0, and

n is an integer from 1 to 4,

or

X represents a bond, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,

Y represents 1,4-piperazinylene,

m is an integer from 1 to 4, and

n is an integer from 1 to 4,

or

A represents -Y-X-C_mH_{2m}-Z-C_nH_{2n}-, wherein

X represents a bond, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,

Y represents 4,1-piperidinylene,

Z represents a bond, -O-, -S-, -S(O)₂- or -C(O)-NH-,

m is an integer from 1 to 4, and

n is an integer from 1 to 4,

or

X represents a bond, -O-, -S-, -C(O)-, -S(O)₂-, -C(O)-NH- or -C(S)-NH-,

Y represents 1,4-phenylene, 1,3-phenylene, 1,4-cyclohexylene, 1,3-cyclohexylene or 1,3-cyclopentylene,

Z represents a bond, -O-, -S-, -S(O)₂- or -C(O)-NH-,

m is an integer from 1 to 4, and

n is an integer from 1 to 4,

R8 is hydrogen, methyl or ethyl,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein

R9 is hydrogen, halogen, hydroxyl, amino, ureido [-NH-C(O)-NH₂], formylamino [-NH-C(O)H], 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,

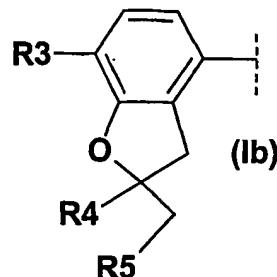
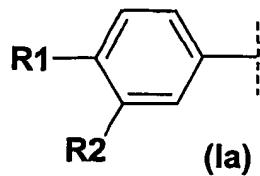
R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl or 1-4C-alkylcarbonyloxy,

R11 is hydrogen or halogen,

and the salts of these compounds.

3. Compounds of formula I according to claim 1 in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



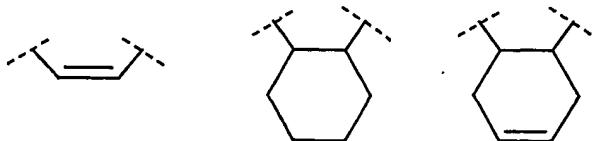
wherein

- R1 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is 1-2C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is methyl and
- R5 is hydrogen,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane or cyclohexane ring,

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



and A either represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein

either

- X represents a bond,
- Y represents a bond,
- m is an integer from 1 to 4, and
- n is an integer from 1 to 4,
- or
- X represents a bond, -O- or -C(O)-NH-,
- Y represents 1,4-phenylene or 1,4-cyclohexylene,

m is an integer from 0 to 4, and
n is an integer from 1 to 4,
or
X represents a bond, -C(O)-, -S(O)₂- or -C(S)-NH-,
Y represents 4,1-piperidinylene,
m is 0, and
n is an integer from 1 to 4,

or

A represents -Y-X-C_mH_{2m}-Z-C_nH_{2n}-, wherein

either
X represents a bond or -C(O)-,
Y represents 4,1-piperidinylene,
Z represents a bond, -S- or -S(O)₂-,
m is an integer from 1 to 4, and
n is an integer from 1 to 4,
or
X represents a bond, -O- or -C(O)-NH-,
Y represents 1,4-phenylene or 1,4-cyclohexylene,
Z represents a bond,
m is an integer from 1 to 4, and
n is an integer from 1 to 4,

R8 is hydrogen,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein

R9 is hydrogen, hydroxyl or amino,

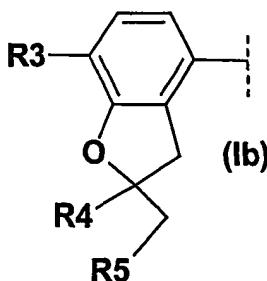
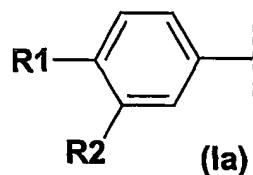
R10 is hydrogen, halogen, cyano, trifluoromethyl or hydroxymethyl,

R11 is hydrogen or halogen,

and the salts of these compounds.

4. Compounds of formula I according to claim 1 in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

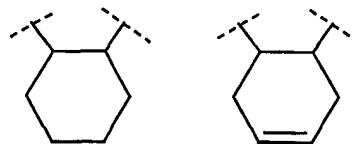
R1 is methoxy or ethoxy,

R2 is methoxy or ethoxy,

R3 is methoxy,

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane ring,

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



and A either represents $-C_mH_{2m}-Y-X-C_nH_{2n}-$, wherein

either

X represents a bond,

Y represents a bond,

m is an integer from 1 to 4, and

n is an integer from 1 to 4,

or

X represents a bond, -O- or -C(O)-NH-,

Y represents 1,4-phenylene,

m is an integer from 0 to 1, and

n is an integer from 1 to 4,

or

X represents a bond, -C(O)-, -S(O)₂- or -C(S)-NH-,

Y represents 4,1-piperidinylene,

m is 0, and

n is an integer from 1 to 4,

or

A represents $-Y-X-C_mH_{2m}-Z-C_nH_{2n}-$, wherein

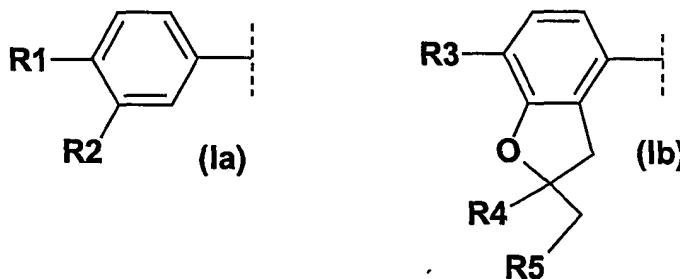
- X represents a bond or $-C(O)-$,
- Y represents 4,1-piperidinylene,
- Z represents a bond, $-S-$ or $-S(O)_2-$,
- m is 2 or 3, and
- n is 2 or 3,

R8 is hydrogen,

Ar₂ is phenyl, 4-amino-3-chloro-5-cyanophenyl, 4-amino-3-chloro-5-trifluoromethylphenyl, 4-hydroxy-3-hydroxymethylphenyl, 4-amino-3-cyanophenyl or 4-amino-3,5-dichlorophenyl,
and the salts of these compounds.

5. Compounds of formula I according to claim 1 in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

R1 is hydroxyl, halogen, 1-4C-alkoxy or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

R2 is hydroxyl, halogen, 1-8C-alkoxy, 3-7C-cycloalkoxy, 3-7C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

R3 is 1-4C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-4C-alkoxy which is completely or predominantly substituted by fluorine,

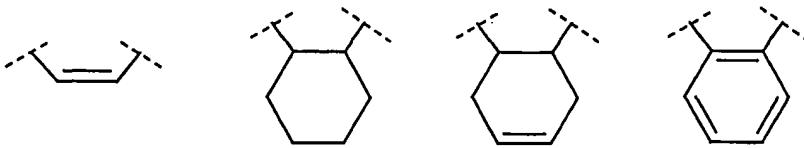
R4 is 1-4C-alkyl and

R5 is hydrogen or 1-4C-alkyl,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked 5-, 6- or 7-membered hydrocarbon ring, optionally interrupted by an oxygen or sulphur atom,

R6 and R7 represent independently from one another hydrogen or 1-4C-alkyl, or R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents $-(CH_2)_m-Y-X-(CH_2)_n-$, wherein

- X represents a bond, -O- (oxygen), -S- (sulfur), -NH-, -C(O)-, -C(O)-NH- or -NH-C(O)-NH-,
- Y represents a bond, phenylene, 4-8C-cycloalkylene or azacycloalkylene,
- m is an integer from 0 to 4,
- n is an integer from 1 to 4,

R8 is hydrogen or 1-4C-alkyl,

Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11,

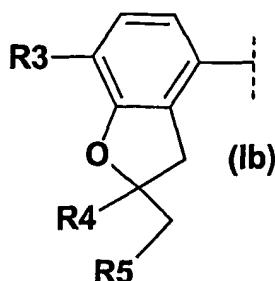
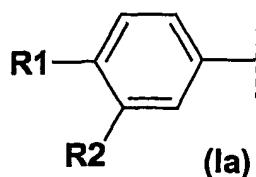
wherein

- R9 is hydrogen, halogen, hydroxyl, amino, ureido [-NH-C(O)-NH₂], formylamino [-NH-C(O)H], 1-4C-alkylcarbonylamino, di-1-4C-alkylaminocarbonyloxy, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl, 1-4C-alkylcarbonyloxy, 1-4C-alkyl, 1-4C-alkoxy, 1-4C-alkylsulfonylamino, 1-4C-alkylsulfonylmethyl or 1-4C-alkoxy-1-4C-alkyl,
- R10 is hydrogen, halogen, hydroxyl, cyano, trifluoromethyl, tolylcarbonyloxy [-O-C(O)-C₆H₄-CH₃], hydroxymethyl or 1-4C-alkylcarbonyloxy,
- R11 is hydrogen or halogen,

and the salts of these compounds.

6. Compounds of formula I according to claim 1 in which

Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



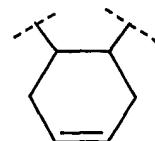
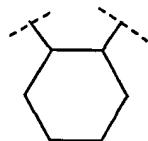
wherein

- R1 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R2 is 1-2C-alkoxy, 3-5C-cycloalkoxy, 3-5C-cycloalkylmethoxy, or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R3 is 1-2C-alkoxy or 1-2C-alkoxy which is completely or predominantly substituted by fluorine,
- R4 is methyl and
- R5 is hydrogen,

or wherein

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane or cyclohexane ring,

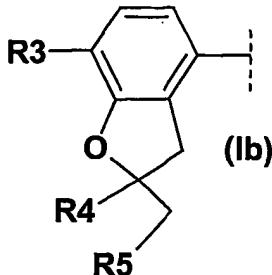
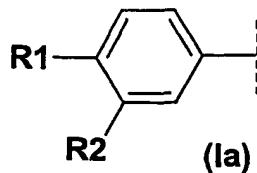
R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



- A represents $-(CH_2)_m-Y-X-(CH_2)_n-$, wherein
 - X represents a bond or $-C(O)-$,
 - Y represents a bond, 1,4-phenylene, 1,4-cyclohexylene or 4,1-piperidinylene,
 - m is an integer from 0 to 4,
 - n is an integer from 1 to 4,
- R8 is hydrogen,
- Ar₂ represents a 8-hydroxy-1H-quinolin-2-on-5-yl radical or a phenyl radical substituted by R9, R10 and R11, wherein
 - R9 is hydrogen, hydroxyl or amino,
 - R10 is hydrogen, halogen, cyano, trifluoromethyl or hydroxymethyl,
 - R11 is hydrogen or halogen,

and the salts of these compounds.

7. Compounds of formula I according to claim 1 in which
 Ar₁ represents a phenyl radical of formula (Ia) substituted by R1 and R2 or a dihydrobenzofuranyl radical of formula (Ib) substituted by R3, R4 and R5



wherein

R1 is methoxy, ethoxy or difluoromethoxy,

R2 is methoxy or ethoxy,

R3 is methoxy,

R4 and R5 together and with inclusion of the two carbon atoms, to which they are bonded, form a spiro-linked cyclopentane ring,

R6 and R7 together and with inclusion of the two carbon atoms, to which they are bonded, form a group selected from



A represents -(CH₂)_m-Y-X-(CH₂)_n-, wherein

X represents a bond,

Y represents a bond or 1,4-phenylene,

m is an integer from 0 to 4,

n is an integer from 1 to 4,

R8 is hydrogen,

Ar₂ is phenyl, 4-amino-3-chloro-5-cyanophenyl, 4-amino-3-chloro-5-trifluoromethyl, 4-hydroxy-3-hydroxymethylphenyl, 4-amino-3-cyanophenyl or 4-amino-3,5-dichlorophenyl,

and the salts of these compounds.

8. Medicaments containing one or more compounds of formula I according to claim 1 together with the usual pharmaceutical auxiliaries and/or carrier materials.
9. Compounds of formula I according to claim 1 for use in the treatment of illnesses.
10. Use of compounds of formula I according to claim 1 for the production of medicaments for the treatment of airway disorders.

INTERNATIONAL SEARCH REPORT

Intern: Application No

PCT/EP 01/06230

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D237/14 C07D237/22 C07D237/32 C07D405/04 A61K31/50
 A61K31/501 A61K31/502 A61P11/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, PAJ, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 99 31090 A (BYK GULDEN LOMBERG CHEM FAB; ULRICH WOLF RUEDIGER (DE); STERK GEER) 24 June 1999 (1999-06-24) cited in the application page 21 -page 26; claims 1,7-9 ---	1,8-10
A	EP 0 934 933 A (BYK GULDEN LOMBERG CHEM FAB) 11 August 1999 (1999-08-11) cited in the application column 8, line 19 -column 10, line 37; claims 1,5-7 ---	1,8-10
A	EP 0 763 534 A (MERCK PATENT GMBH) 19 March 1997 (1997-03-19) cited in the application page 7, line 11 - line 26; claims 1,6-10 --- -/-	1,8-10

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the International filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the International filing date but later than the priority date claimed

- *T* later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the International search

19 July 2001

Date of mailing of the International search report

03/08/2001

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
 Fax (+31-70) 340-3016

Authorized officer

Hass, C

INTERNATIONAL SEARCH REPORT

Inter	Application No
PCT/EP 01/06230	

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 99 31071 A (BYK GULDEN LOMBERG CHEM FAB; STERK GEERT JAN (NL)) 24 June 1999 (1999-06-24) cited in the application page 21 -page 24; claims 1,8-10 ---	1,8-10
A	WO 98 31674 A (BYK GULDEN LOMBERG CHEM FAB; STERK GEERT JAN (NL)) 23 July 1998 (1998-07-23) cited in the application page 19 -page 40; claims 1,8-10 ---	1,8-10
A	EP 0 030 835 A (MORISHITA PHARMA) 24 June 1981 (1981-06-24) claims 1,24 ---	1,8,9
A	PATENT ABSTRACTS OF JAPAN vol. 007, no. 263 (C-196), 24 November 1983 (1983-11-24) & JP 58 146571 A (MORISHITA SEIYAKU KK), 1 September 1983 (1983-09-01) abstract ---	1,8,9
A	PL 164 079 B (AKAD MEDYCZNA) 30 June 1994 (1994-06-30) abstract; figure 1 ---	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

Inten Application No

PCT/EP 01/06230

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9931090	A 24-06-1999	AU 2270199 A		05-07-1999
		EP 1042319 A		11-10-2000
		PL 341239 A		26-03-2001
EP 0934933	A 11-08-1999	NONE		
EP 0763534	A 19-03-1997	DE 19533975 A		20-03-1997
		AU 716113 B		17-02-2000
		AU 6551796 A		20-03-1997
		BR 9603736 A		26-05-1998
		CA 2185397 A		15-03-1997
		CN 1157287 A		20-08-1997
		CZ 9602630 A		18-03-1998
		HU 9602511 A		28-03-1997
		JP 9124611 A		13-05-1997
		NO 963852 A		17-03-1997
		PL 316070 A		17-03-1997
		SK 110096 A		06-08-1997
		US 5859008 A		12-01-1999
		ZA 9607766 A		26-03-1997
WO 9931071	A 24-06-1999	AU 1760399 A		05-07-1999
WO 9831674	A 23-07-1998	AU 5862998 A		07-08-1998
		BR 9806752 A		14-03-2000
		CN 1249749 T		05-04-2000
		EP 0971901 A		19-01-2000
		HU 0001541 A		28-05-2001
		NO 993301 A		10-09-1999
		PL 334561 A		13-03-2000
		SK 95199 A		10-12-1999
		TR 9901653 T		21-10-1999
		US 6103718 A		15-08-2000
EP 0030835	A 24-06-1981	JP 1491255 C		07-04-1989
		JP 57028060 A		15-02-1982
		JP 63038993 B		03-08-1988
		JP 57028061 A		15-02-1982
		JP 57028054 A		15-02-1982
		JP 1480814 C		10-02-1989
		JP 57028055 A		15-02-1982
		JP 63025587 B		26-05-1988
		JP 57028056 A		15-02-1982
		JP 1435448 C		07-04-1988
		JP 56081568 A		03-07-1981
		JP 62042900 B		10-09-1987
		AT 3288 T		15-05-1983
		DE 3063196 D		16-06-1983
		ES 498263 D		16-12-1981
		ES 8201553 A		16-03-1982
		ZA 8007209 A		25-11-1981
JP 58146571	A 01-09-1983	NONE		
PL 164079	B 30-06-1994	NONE		